

**PROTEIN-PROTEIN INTERACTIONS
IN ADIPOCYTE CELLS**



CROSS REFERENCE TO RELATED APPLICATIONS

This application claims priority under 35 U.S.C. § 119(e) to U.S. Provisional Application No. 60/259,377, filed January 2, 2001.

BACKGROUND OF THE INVENTION

Most biological processes involve specific protein-protein interactions. Protein-protein interactions enable two or more proteins to associate. A large number of non-covalent bonds form between the proteins when two protein surfaces are precisely matched. These bonds account for the specificity of recognition. Thus, protein-protein interactions are involved, for example, in the assembly of enzyme subunits, in antibody-antigen recognition, in the formation of biochemical complexes, in the correct folding of proteins, in the metabolism of proteins, in the transport of proteins, in the localization of proteins, in protein turnover, in first translation modifications, in the core structures of viruses and in signal transduction.

General methodologies to identify interacting proteins or to study these interactions have been developed. Among these methods are the two-hybrid system originally developed by Fields and co-workers and described, for example, in U.S. Patent Nos. 5,283,173, 5,468,614 and 5,667,973, which are hereby incorporated by reference.

The earliest and simplest two-hybrid system, which acted as basis for development of other versions, is an *in vivo* assay between two specifically constructed proteins. The first protein, known in the art as the "bait protein" is a chimeric protein which binds to a site on DNA upstream of a reporter gene by means of a DNA-binding domain or BD. Commonly, the binding domain is the DNA-binding domain from either Gal4 or native *E. coli* LexA and the sites placed upstream of the reporter are Gal4 binding sites or LexA operators, respectively.

fact, a doctrine has been developed in this field for interpreting the results and in common practice an additional technique such as co-immunoprecipitation or gradient sedimentation of the putative interactors from the appropriate cell or tissue type are generally performed. The methods used for interpreting the results are described by Brent and Finley, Jr. in *Ann. Rev. Genet.*, 31 pgs. 663-704 (1997). Thus, the data interpretation is very questionable using the conventional systems.

One method to overcome the difficulties encountered with the methods in the prior art is described in WO99/42612, incorporated herein by reference. This method is similar to the two-hybrid system described in the prior art in that it also uses bait and prey polypeptides. However, the difference with this method is that a step of mating at least one first haploid recombinant yeast cell containing the prey polypeptide to be assayed with a second haploid recombinant yeast cell containing the bait polynucleotide is performed. Of course the person skilled in the art would appreciate that either the first recombinant yeast cell or the second recombinant yeast cell also contains at least one detectable reporter gene that is activated by a polypeptide including a transcriptional activation domain.

The method described in WO99/42612 permits the screening of more prey polynucleotides with a given bait polynucleotide in a single step than in the prior art systems due to the cell to cell mating strategy between haploid yeast cells. Furthermore, this method is more thorough and reproducible, as well as sensitive. Thus, the presence of false negatives and/or false positives is extremely minimal as compared to the conventional prior art methods.

The causes of Non-insulin dependent diabetes mellitus (NIDDM) and obesity are often related to defects or problems with adipose tissue. Adipocytes play a critical role in lipid storage and metabolism. Adipocytes also act as endocrine cells to influence physiological parameters such as insulin sensitivity and body weight (Flier, et al., *Cell*, (1995) 80: 15-18). For example, the *ob* gene encodes leptin, an adipocyte secreted endocrine factor (Zhang, et al., *Nature* (1994) 372: 425-432). Leptin has been shown to reduce body weight and blood glucose in obese, diabetic rodents (Pellemounter, et al., *Science*, (1995) 269: 540-543).

NIDDM is treated predominately with insulin. However, insulin is not convenient to use in that it must be injected 2-4 times per day and must be stored properly to prevent

loss of efficacy. Other drugs used to treat NIDDM include troglitazone ("Rezulin"), a PPAR γ agonist, Glucophage and sulfonylureas. Unfortunately, there are safety concerns related to the use of these drugs. The identification of safe, effective, orally available drugs for the treatment of NIDDM would greatly enhance the quality of life of patients who suffer from this disease.

Several adipocyte-specific enzymes and receptors have been shown to be important targets for anti-obesity and anti-diabetic drug discovery. For example, agonists of the β_3 adrenergic receptor, which is found predominantly in the adipose tissue in man (Arner, et al., New England Journal of Medicine, (1995) 333: 382-383), have anti-obesity and anti-diabetic properties in rodents and are currently in phase II/III trials in man. The thiazolidinedione class of compounds (TZDs), including troglitazone and ciglitazone, has been shown to improve insulin sensitivity and thereby reduce hyperglycemia and hyperlipidemia conditions in rodents and in humans (Saltiel, et al., Diabetes, (1996) 45: 1661-1669; Sreenan, et al., American Journal Physiol, (1996) 271: E742-E747; Nolan, et al., New England Journal of Medicine, (1994) 331: 1188-1193. Troglitazone (Rezulin") is approved for use in the U. S. and Japan. Many TZDs, including troglitazone and ciglitazone, are potent activators of Peroxisome Proliferator Activated Receptor gamma (PPAR γ), a member of the nuclear receptor family of transcription factors (Tontonoz, et al., Cell, (1994) 79: 1147-1156; Lehmann, et al., Journal of Biological Chemistry, (1995) 270: 12953-12955). PPAR β , is a key regulator of adipocyte differentiation and is most abundant in adipose tissue.

This shows that it is still needed to explore all mechanisms of adipocyte differentiation and to identify drug targets for metabolism diseases.

SUMMARY OF THE INVENTION

Thus, it is an object of the present invention to identify protein-protein interactions in adipocytes.

It is another object of the present invention to identify protein-protein interactions in adipocytes for the development of more effective and better targeted therapeutic applications.

It is yet another object of the present invention to identify complexes of polypeptides or polynucleotides encoding the polypeptides and fragments of the polypeptides of adipocytes.

It is yet another object of the present invention to identify antibodies to these complexes of polypeptides or polynucleotides encoding the polypeptides and fragments of the polypeptides of adipocytes including polyclonal, as well as monoclonal antibodies that are used for detection.

It is still another object of the present invention to identify selected interacting domains of the polypeptides, called SID® polypeptides.

It is still another object of the present invention to identify selected interacting domains of the polynucleotides, called SID® polynucleotides.

It is another object of the present invention to generate protein-protein interactions maps called PIM®s.

It is yet another object of the present invention to provide a method for screening drugs for agents which modulate the interaction of proteins and pharmaceutical compositions that are capable of modulating the protein-protein interactions in adipocytes.

It is another object to administer the nucleic acids of the present invention via gene therapy.

It is yet another object of the present invention to provide protein chips or protein microarrays.

It is yet another object of the present invention to provide a report in, for example paper, electronic and/or digital forms, concerning the protein-protein interactions, the modulating compounds and the like as well as a PIM®.

These and other objects are achieved by the present invention as evidenced by the following paragraphs, detailed description of the invention and the claims.

Thus the present invention relates to a complex of interacting proteins of columns 1 and 3 of Table 2.

Furthermore, the present invention provides SID® polynucleotides and SID® polypeptides, as well as a PIM® for adipocytes.

Furthermore, the present invention provides scientific evidence of protein interactions between MT1R and MUPP1, as well as between β arrestin2 and Oct-1 have been confirmed in adipocytes.

The present invention also provides antibodies to the protein-protein complexes in adipocytes.

In another aspect, the present invention relates to the interaction between the MT1A receptor with MUPP1. Melatonin (the hormone of darkness) is involved in the regulation of circadian rhythms and sleep, but it also has roles in visual, cerebrovascular, reproductive, neuroendocrine, and neuroimmunological functions. Melatonin mediates its effects through G protein-coupled receptors (GPCR): MT(1), MT(2), and, possibly, MT(3). Information is provided about the interaction of MT1A receptor with MUPP1, a 13 PDZ domains containing molecule. MUPP1 which has previously been shown to interact with the 5-HT(2C) serotonin receptor may serve as a multivalent scaffold protein that selectively assembles and targets signaling complexes to the MT1A receptor and therefore may modulate its activity and consequently the physiological roles attributed to this receptor.

In the classical model of G-protein-coupled receptor (GPCR) regulation, arrestins terminate receptor signalling. More recently, arrestins have been shown to link GPCRs to several signalling pathways, including activation of the non-receptor tyrosine kinase SRC and mitogen-activated protein kinase. In these cascades, arrestins function as adaptors and scaffolds, bringing sequentially acting kinases into proximity with each other and the receptor. Here, we provide evidences for an interaction between beta-arrestin 2 and Oct-1, a ubiquitously expressed member of the POU family of transcription factor which is involved in the regulation of a wide variety of genes

implicated in cell cycle regulation, development and hormonal signals. Moreover, we have shown that beta arrestin 2 binding to Oct-1 modulate its transcriptional activity. These data indicate that GPCR signaling may modulate through arrestin the activity of this class of important transcription factors.

Another aspect of the present invention provides a method for screening drugs for agents that modulate the protein-protein interactions and pharmaceutical compositions that are capable of modulating protein-protein interactions.

Yet another aspect of the present invention provides protein chips or protein microarrays.

Yet another aspect of the present invention provides a report in, for example, paper, electronic and/or digital forms.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a schematic representation of the pB1 plasmid.

Fig. 2 is a schematic representation of the pB5 plasmid.

Fig. 3 is a schematic representation of the pB6 plasmid.

Fig. 4 is a schematic representation of the pB13 plasmid.

Fig. 5 is a schematic representation of the pB14 plasmid.

Fig. 6 is a schematic representation of the pB20 plasmid.

Fig. 7 is a schematic representation of the pP1 plasmid.

Fig. 8 is a schematic representation of the pP2 plasmid.

Fig. 9 is a schematic representation of the pP3 plasmid.

Fig. 10 is a schematic representation of the pP6 plasmid.

Fig. 11 is a schematic representation of the pP7 plasmid.

Fig. 12 is a schematic representation of vectors expressing the T25 fragment.

Fig. 13 is a schematic representation of vectors expressing the T18 fragment.

Fig. 14 is a schematic representation of various vectors of pCmAHL1, pT25 and pT18.

Fig. 15 is a schematic representation identifying the SID®s of adipocytes. In this figure the "Full-length prey protein" is the Open Reading Frame (ORF) or coding sequence (CDS) where the identified prey polypeptides are included. The Selected

Interaction Domain (SID®) is determined by the commonly shared polypeptide domain of every selected prey fragment.

Fig. 16 is a protein map (PIM®).

Fig. 17 are Western blots verifying the interaction between MTR1 (melatonin 1 receptors) and MUPP1(multi-PDZ-domain protein) in whole cell lysates of HEK 293 cells transfected with both cDNAs. Flag-tagged MT1 receptors were immunoprecipitated with anti Flag antibodies and MUPP1 was detected with an anti-MUPP1 antibody.

Fig. 18 is a graph also verifying the interaction between MTR1 and MUPP1 in BRET experiments. Expression of MUPP1 decreased the energy transfer between MT1R-Rluc and MTR1-YFP in a dose dependent manner (Fig. 18B). The transfer between MT2R-Rluc and MTR2-YFP was insensitive to MUPP1 expression confirming the specificity of the interaction (Fig. 18A).

Fig. 19 is a graph illustrating that β arrestin2 has an inhibitory effect on Oct-1-mediated gene expression.(Octamer binding protein-1).

DETAILED DESCRIPTION OF THE INVENTION

The adipocytes (undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes) studied in the present invention are obtained by the method described in the PCT patent application WO96/34100.

As used herein the terms "polynucleotides", "nucleic acids" and "oligonucleotides" are used interchangeably and include, but are not limited to RNA, DNA, RNA/DNA sequences of more than one nucleotide in either single chain or duplex form. The polynucleotide sequences of the present invention may be prepared from any known method including, but not limited to, any synthetic method, any recombinant method, any *ex vivo* generation method and the like, as well as combinations thereof.

The term "polypeptide" means herein a polymer of amino acids having no specific length. Thus, peptides, oligopeptides and proteins are included in the definition of "polypeptide" and these terms are used interchangeably throughout the specification, as well as in the claims. The term "polypeptide" does not exclude post-translational modifications such as polypeptides having covalent attachment of glycosyl groups, acetyl groups, phosphate groups, lipid groups and the like. Also encompassed by this definition of "polypeptide" are homologs thereof.

By the term "homologs" is meant structurally similar genes contained within a given species, orthologs are functionally equivalent genes from a given species or strain, as determined for example, in a standard complementation assay. Thus, a polypeptide of interest can be used not only as a model for identifying similar genes in given strains, but also to identify homologs and orthologs of the polypeptide of interest in other species. The orthologs, for example, can also be identified in a conventional complementation assay. In addition or alternatively, such orthologs can be expected to exist in bacteria (or other kind of cells) in the same branch of the phylogenetic tree, as set forth, for example, at <ftp://ftp.cme.msu.edu/pub/rdp/SSU-rRNA/SSU/Prok.phylo>.

As used herein the term "prey polynucleotide" means a chimeric polynucleotide encoding a polypeptide comprising (i) a specific domain; and (ii) a polypeptide that is to be tested for interaction with a bait polypeptide. The specific domain is preferably a transcriptional activating domain.

As used herein, a "bait polynucleotide" is a chimeric polynucleotide encoding a chimeric polypeptide comprising (i) a complementary domain; and (ii) a polypeptide that is to be tested for interaction with at least one prey polypeptide. The complementary domain is preferably a DNA-binding domain that recognizes a binding site that is further detected and is contained in the host organism.

As used herein "complementary domain" is meant a functional constitution of the activity when bait and prey are interacting; for example, enzymatic activity.

As used herein "specific domain" is meant a functional interacting activation domain that may work through different mechanisms by interacting directly or indirectly through intermediary proteins with RNA polymerase II or III-associated proteins in the vicinity of the transcription start site.

As used herein the term "complementary" means that, for example, each base of a first polynucleotide is paired with the complementary base of a second polynucleotide whose orientation is reversed. The complementary bases are A and T (or A and U) or C and G.

The term "sequence identity" refers to the identity between two peptides or between two nucleic acids. Identity between sequences can be determined by comparing a position in each of the sequences which may be aligned for the purposes of comparison. When a position in the compared sequences is occupied by the same base or amino acid, then the sequences are identical at that position. A degree of sequence identity between nucleic acid sequences is a function of the number of identical nucleotides at positions shared by these sequences. A degree of identity between amino acid sequences is a function of the number of identical amino acid sequences that are shared between these sequences. Since two polypeptides may each (i) comprise a sequence (i.e., a portion of a complete polynucleotide sequence) that is similar between two polynucleotides, and (ii) may further comprise a sequence that is divergent between two polynucleotides, sequence identity comparisons between two or more polynucleotides over a "comparison window" refers to the conceptual segment of at least 20 contiguous nucleotide positions wherein a polynucleotide sequence may be compared to a reference nucleotide sequence of at least 20 contiguous nucleotides and wherein the portion of the polynucleotide sequence in the comparison window may comprise additions or deletions (i.e., gaps) of 20 percent or less compared to the reference sequence (which does not comprise additions or deletions) for optimal alignment of the two sequences.

To determine the percent identity of two amino acids sequences or two nucleic acid sequences, the sequences are aligned for optimal comparison. For example, gaps can be introduced in the sequence of a first amino acid sequence or a first nucleic acid sequence for optimal alignment with the second amino acid sequence or second nucleic acid sequence. The amino acid residues or nucleotides at corresponding amino acid positions or nucleotide positions are then compared. When a position in the first sequence is occupied by the same amino acid residue or nucleotide as the corresponding position in the second sequence, the molecules are identical at that position.

The percent identity between the two sequences is a function of the number of identical positions shared by the sequences. Hence % identity = number of identical positions / total number of overlapping positions X 100.

In this comparison the sequences can be the same length or may be different in length. Optimal alignment of sequences for determining a comparison window may be conducted by the local homology algorithm of Smith and Waterman (*J. Theor. Biol.*, 91 (2) pgs. 370-380 (1981), by the homology alignment algorithm of Needleman and Wunsch, *J.*

Mol. Biol., 48(3) pgs. 443-453 (1972), by the search for similarity via the method of Pearson and Lipman, *PNAS, USA*, 85(5) pgs. 2444-2448 (1988) , by computerized implementations of these algorithms (GAP, BESTFIT, FASTA and TFASTA in the Wisconsin Genetics Software Package Release 7.0, Genetic Computer Group, 575, Science Drive, Madison, Wisconsin) or by inspection.

The best alignment (i.e., resulting in the highest percentage of identity over the comparison window) generated by the various methods is selected.

The term "sequence identity" means that two polynucleotide sequences are identical (i.e., on a nucleotide by nucleotide basis) over the window of comparison. The term "percentage of sequence identity" is calculated by comparing two optimally aligned sequences over the window of comparison, determining the number of positions at which the identical nucleic acid base (e.g., A, T, C, G, U, or I) occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the window of comparison (i.e., the window size) and multiplying the result by 100 to yield the percentage of sequence identity. The same process can be applied to polypeptide sequences.

The percentage of sequence identity of a nucleic acid sequence or an amino acid sequence can also be calculated using BLAST software (Version 2.06 of September 1998) with the default or user defined parameter.

The term "sequence similarity" means that amino acids can be modified while retaining the same function. It is known that amino acids are classified according to the nature of their side groups and some amino acids such as the basic amino acids can be interchanged for one another while their basic function is maintained.

The term "isolated" as used herein means that a biological material such as a nucleic acid or protein has been removed from its original environment in which it is naturally present. For example, a polynucleotide present in a plant, mammal or animal is present in its natural state and is not considered to be isolated. The same polynucleotide separated from the adjacent nucleic acid sequences in which it is naturally inserted in the genome of the plant or animal is considered as being "isolated."

The term "isolated" is not meant to exclude artificial or synthetic mixtures with other compounds, or the presence of impurities which do not interfere with the biological activity and which may be present, for example, due to incomplete purification, addition of stabilizers or mixtures with pharmaceutically acceptable excipients and the like.

"Isolated polypeptide" or "isolated protein" as used herein means a polypeptide or protein which is substantially free of those compounds that are normally associated with the polypeptide or protein in a naturally state such as other proteins or polypeptides, nucleic acids, carbohydrates, lipids and the like.

The term "purified" as used herein means at least one order of magnitude of purification is achieved, preferably two or three orders of magnitude, most preferably four or five orders of magnitude of purification of the starting material or of the natural material. Thus, the term "purified" as utilized herein does not mean that the material is 100% purified and thus excludes any other material.

The term "variants" when referring to, for example, polynucleotides encoding a polypeptide variant of a given reference polypeptide are polynucleotides that differ from the reference polypeptide but generally maintain their functional characteristics of the reference polypeptide. A variant of a polynucleotide may be a naturally occurring allelic variant or it may be a variant that is known naturally not to occur. Such non-naturally occurring variants of the reference polynucleotide can be made by, for example, mutagenesis techniques, including those mutagenesis techniques that are applied to polynucleotides, cells or organisms.

Generally, differences are limited so that the nucleotide sequences of the reference and variant are closely similar overall and, in many regions identical.

Variants of polynucleotides according to the present invention include, but are not limited to, nucleotide sequences which are at least 95% identical after alignment to the reference polynucleotide encoding the reference polypeptide. These variants can also have 96%, 97%, 98% and 99.999% sequence identity to the reference polynucleotide.

Nucleotide changes present in a variant polynucleotide may be silent, which means that these changes do not alter the amino acid sequences encoded by the reference polynucleotide.

Substitutions, additions and/or deletions can involve one or more nucleic acids. Alterations can produce conservative or non-conservative amino acid substitutions, deletions and/or additions.

Variants of a prey or a SID® polypeptide encoded by a variant polynucleotide can possess a higher affinity of binding and/or a higher specificity of binding to its protein or polypeptide counterpart, against which it has been initially selected. In another context, variants can also lose their ability to bind to their protein or polypeptide counterpart.

By "anabolic pathway" is meant a reaction or series of reactions in a metabolic pathway that synthesize complex molecules from simpler ones, usually requiring the input of energy. An anabolic pathway is the opposite of a catabolic pathway.

As used herein, a "catabolic pathway" is a series of reactions in a metabolic pathway that break down complex compounds into simpler ones, usually releasing energy in the process. A catabolic pathway is the opposite of an anabolic pathway.

As used herein, "drug metabolism" is meant the study of how drugs are processed and broken down by the body. Drug metabolism can involve the study of enzymes that break down drugs, the study of how different drugs interact within the body and how diet and other ingested compounds affect the way the body processes drugs.

As used herein, "metabolism" means the sum of all of the enzyme-catalyzed reactions in living cells that transform organic molecules.

By "secondary metabolism" is meant pathways producing specialized metabolic products that are not found in every cell.

As used herein, "SID®" means a Selected Interacting Domain and is identified as follows: for each bait polypeptide screened, selected prey polypeptides are compared. Overlapping fragments in the same ORF or CDS define the selected interacting domain.

As used herein the term "PIM®" means a protein-protein interaction map. This map is obtained from data acquired from a number of separate screens using different bait polypeptides and is designed to map out all of the interactions between the polypeptides.

The term "affinity of binding", as used herein, can be defined as the affinity constant K_a when a given SID® polypeptide of the present invention which binds to a polypeptide and is the following mathematical relationship:

$$K_a = \frac{[\text{SID®/polypeptide complex}]}{[\text{free SID®}] [\text{free polypeptide}]}$$

wherein [free SID®], [free polypeptide] and [SID®/polypeptide complex] consist of the concentrations at equilibrium respectively of the free SID® polypeptide, of the free polypeptide onto which the SID® polypeptide binds and of the complex formed between SID® polypeptide and the polypeptide onto which said SID® polypeptide specifically binds.

The affinity of a SID® polypeptide of the present invention or a variant thereof for its polypeptide counterpart can be assessed, for example, on a Biacore™ apparatus marketed by Amersham Pharmacia Biotech Company such as described by Szabo et al *Curr Opin Struct Biol* 5 pgs. 699-705 (1995) and by Edwards and Leartherbarrow, *Anal. Biochem* 246 pgs. 1-6 (1997).

As used herein the phrase "at least the same affinity" with respect to the binding affinity between a SID® polypeptide of the present invention to another polypeptide means that the K_a is identical or can be at least two-fold, at least three-fold or at least five fold greater than the K_a value of reference.

As used herein, the term "modulating compound" means a compound that inhibits or stimulates or can act on another protein which can inhibit or stimulate the protein-protein interaction of a complex of two polypeptides or the protein-protein interaction of two polypeptides.

More specifically, the present invention comprises complexes of polypeptides or polynucleotides encoding the polypeptides composed of a bait polypeptide, or a bait polynucleotide encoding a bait polypeptide and a prey polypeptide or a prey polynucleotide encoding a prey polypeptide. The prey polypeptide or prey polynucleotide

encoding the prey polypeptide is capable of interacting with a bait polypeptide of interest in various hybrid systems.

As described in the Background of the present invention there are various methods known in the art to identify prey polypeptides that interact with bait polypeptides of interest. These methods, include, but are not limited to, generic two-hybrid systems as described by Fields et al in *Nature*, 340:245-246 (1989) and more specifically in U.S. Patent Nos. 5,283,173, 5,468,614 and 5,667,973, which are hereby incorporated by reference; the reverse two-hybrid system described by Vidal et al, *supra*; the two plus one hybrid method described, for example, in Tirode et al, *supra*; the yeast forward and reverse 'n'-hybrid systems as described in Vidal and Legrain, *supra*; the method described in WO 99/42612; those methods described in Legrain et al *FEBS Letters* 480 pgs. 32-36 (2000) and the like.

The present invention is not limited to the type of method utilized to detect protein-protein interactions and therefore any method known in the art and variants thereof can be used. It is however better to use the method described in WO99/42612 or WO00/66722, both references incorporated herein by reference due to the methods' sensitivity, reproducibility and reliability.

Protein-protein interactions can also be detected using complementation assays such as those described by Pelletier et al at <http://www.abrf.org/JBT/Articles/JBT0012/jbt0012.html>, WO 00/07038 and WO98/34120.

Although the above methods are described for applications in the yeast system, the present invention is not limited to detecting protein-protein interactions using yeast, but also includes similar methods that can be used in detecting protein-protein interactions in, for example, mammalian systems as described, for example in Takacs et al., *Proc. Natl. Acad. Sci., USA*, **90** (21):10375-79 (1993) and Vasavada et al., *Proc. Natl. Acad. Sci., USA*, **88** (23):10686-90 (1991), as well as a bacterial two-hybrid system as described in Karimova et al (1998), WO99/28746, WO 00/66722 and Legrain et al *FEBS Letters*, **480** pgs. 32-36 (2000).

The above-described methods are limited to the use of yeast, mammalian cells and *Escherichia coli* cells, the present invention is not limited in this manner. Consequently, mammalian and typically human cells, as well as bacterial, yeast, fungus, insect,

nematode and plant cells are encompassed by the present invention and may be transfected by the nucleic acid or recombinant vector as defined herein.

Examples of suitable cells include, but are not limited to, VERO cells, HELA cells such as ATCC No. CCL2, CHO cell lines such as ATCC No. CCL61, COS cells such as COS-7 cells and ATCC No. CRL 1650 cells, W138, BHK, HepG2, 3T3 such as ATCC No. CRL6361, A549, PC12, K562 cells, 293 cells, Sf9 cells such as ATCC No. CRL1711 and Cv1 cells such as ATCC No. CCL70.

Other suitable cells that can be used in the present invention include, but are not limited to, prokaryotic host cells strains such as *Escherichia coli*, (e.g., strain DH5- α), *Bacillus subtilis*, *Salmonella typhimurium*, or strains of the genera of *Pseudomonas*, *Streptomyces* and *Staphylococcus*.

Further suitable cells that can be used in the present invention include yeast cells such as those of *Saccharomyces* such as *Saccharomyces cerevisiae*.

The bait polynucleotide, as well as the prey polynucleotide can be prepared according to the methods known in the art such as those described above in the publications and patents reciting the known method *per se*.

The bait polynucleotide of the present invention is obtained from adipocyte's cDNA. The prey polynucleotide is cDNA fragment from a either library of human placenta or undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes, or variants of cDNA fragment from a either library of human placenta or undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes, and fragments from the genome or transcriptome of human placenta or undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes ranging from about 12 to about 5,000, or about 12 to about 10,000 or from about 12 to about 20,000. The prey polynucleotide is then selected, sequenced and identified.

A human placenta or undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes prey library is prepared from the human placenta or undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes, respectively, and constructed in the specially designed prey vector pP6 as shown in Figure 10 after ligation of suitable linkers such that every cDNA insert is fused to a nucleotide sequence in the vector that encodes the transcription activation domain of a reporter gene. Any transcription activation

domain can be used in the present invention. Examples include, but are not limited to, Gal4, YP16, B42, His and the like. Toxic reporter genes, such as CAT^R, CYH2, CYH1, URA3, bacterial and fungi toxins and the like can be used in reverse two-hybrid systems.

The polypeptides encoded by the nucleotide inserts of the human placenta or undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes prey library thus prepared are termed "prey polypeptides" in the context of the presently described selection method of the prey polynucleotides.

The bait polynucleotides can be inserted in bait plasmid pB6 as illustrated in Figure 3. The bait polynucleotide insert is fused to a polynucleotide encoding the binding domain of, for example, the Gal4 DNA binding domain and the shuttle expression vector is used to transform cells.

The bait polynucleotides (column 2.1 of Table 1) and polypeptides (column 2.2 of Table 1) used in the present invention are described in Table 1.

As stated above, any cells can be utilized in transforming the bait and prey polynucleotides of the present invention including mammalian cells, bacterial cells, yeast cells, insect cells and the like.

In an embodiment, the present invention identifies protein-protein interactions in yeast. In using known methods a prey positive clone is identified containing a vector which comprises a nucleic acid insert encoding a prey polypeptide which binds to a bait polypeptide of interest. The method in which protein-protein interactions are identified comprises the following steps:

i) mating at least one first haploid recombinant yeast cell clone from a recombinant yeast cell clone library that has been transformed with a plasmid containing the prey polynucleotide to be assayed with a second haploid recombinant yeast cell clone transformed with a plasmid containing a bait polynucleotide encoding for the bait polypeptide;

ii) cultivating diploid cell clones obtained in step i) on a selective medium; and

iii) selecting recombinant cell clones which grow on the selective medium.

This method may further comprise the step of:

iv) characterizing the prey polynucleotide contained in each recombinant cell clone which is selected in step iii).

In yet another embodiment of the present invention, *in lieu* of yeast, *Escherichia coli* is used in a bacterial two-hybrid system, which encompasses a similar principle to that described above for yeast, but does not involve mating for characterizing the prey polynucleotide.

In yet another embodiment of the present invention, mammalian cells and a method similar to that described above for yeast for characterizing the prey polynucleotide are used.

By performing the yeast, bacterial or mammalian two-hybrid system it is possible to identify for one particular bait an interacting prey polypeptide. The prey polypeptide that has been selected by testing the library of preys in a screen using the two-hybrid, two plus one hybrid methods and the like, encodes the polypeptide interacting with the protein of interest.

The present invention is also directed, in a general aspect, to a complex of polypeptides, polynucleotides encoding the polypeptides composed of a bait polypeptide or bait polynucleotide encoding the bait polypeptide and a prey polypeptide or prey polynucleotide encoding the prey polypeptide capable of interacting with the bait polypeptide of interest. These complexes are identified in Table 2, as the bait amino acid sequences and the prey amino acid sequences, as well as the bait and prey nucleic acid sequences.

In another aspect, the present invention relates to a complex of polynucleotides consisting of a first polynucleotide, or a fragment thereof, encoding a prey polypeptide that interacts with a bait polypeptide and a second polynucleotide or a fragment thereof. This fragment has at least 12 consecutive nucleotides, but can have between 12 and 5,000 consecutive nucleotides, or between 12 and 10,000 consecutive nucleotides or between 12 and 20,000 consecutive nucleotides.

The complexes of the two polypeptides of columns 1 and 3 of Table 2 and the sets of two polynucleotides encoding these polypeptides also form part of the present invention.

In yet another embodiment, the present invention relates to an isolated complex of at least two polypeptides encoded by two polynucleotides wherein said two polypeptides

are associated in the complex by affinity binding and are depicted in columns 1 and 3 of Table 1.

In yet another embodiment, the present invention relates to an isolated complex comprising at least a polypeptide as described in column 1 of Table 2 and a polypeptide as described in column 3 of Table 2. The present invention is not limited to these polypeptide complexes alone but also includes the isolated complex of the two polypeptides in which fragments and/or homologous polypeptides exhibiting at least 95% sequence identity, as well as from 96% sequence identity to 99.999% sequence identity.

Also encompassed in another embodiment of the present invention is an isolated complex in which the SID® of the prey polypeptides encoded by SEQ ID Nos. [15, 16, 17 etc.] in Table 2 forming the isolated complex.

Besides the isolated complexes described above, nucleic acids coding for a Selected Interacting Domain (SID®) polypeptide or a variant thereof or any of the nucleic acids set forth in Table 2 can be inserted into an expression vector which contains the necessary elements for the transcription and translation of the inserted protein-coding sequence. Such transcription elements include a regulatory region and a promoter. Thus, the nucleic acid which may encode a marker compound of the present invention is operably linked to a promoter in the expression vector. The expression vector may also include a replication origin.

A wide variety of host/expression vector combinations are employed in expressing the nucleic acids of the present invention. Useful expression vectors that can be used include, for example, segments of chromosomal, non-chromosomal and synthetic DNA sequences. Suitable vectors include, but are not limited to, derivatives of SV40 and pcDNA and known bacterial plasmids such as col EI, pCR1, pBR322, pMal-C2, pET, pGEX as described by Smith et al [need cite 1988], pMB9 and derivatives thereof, plasmids such as RP4, phage DNAs such as the numerous derivatives of phage I such as NM989, as well as other phage DNA such as M13 and filamentous single stranded phage DNA; yeast plasmids such as the 2 micron plasmid or derivatives of the 2m plasmid, as well as centomeric and integrative yeast shuttle vectors; vectors useful in eukaryotic cells such as vectors useful in insect or mammalian cells; vectors derived from combinations of plasmids and phage DNAs, such as plasmids that have been modified to employ phage DNA or the expression control sequences; and the like.

For example in a baculovirus expression system, both non-fusion transfer vectors, such as, but not limited to pVL941 (*Bam*HI cloning site Summers, pVL1393 (*Bam*HI, *Sma*I, *Xba*I, *Eco*RI, *Not*I, *Xma*III, *Bgl*II and *Pst*I cloning sites; Invitrogen) pVL1392 (*Bgl*II, *Pst*I, *Not*I, *Xma*III, *Eco*RI, *Xba*I, *Sma*I and *Bam*HI cloning site; Summers and Invitrogen) and pBlueBacIII (*Bam*HI, *Bgl*II, *Pst*I, *Nco*I and *Hind*III cloning site, with blue/white recombinant screening, Invitrogen), and fusion transfer vectors such as, but not limited to, pAc700(*Bam*HI and *Kpn*I cloning sites, in which the *Bam*HI recognition site begins with the initiation codon; Summers), pAc701 and pAc70-2 (same as pAc700, with different reading frames), pAc360 (*Bam*HI cloning site 36 base pairs downstream of a polyhedrin initiation codon; Invitrogen (195)) and pBlueBacHisA, B, C (three different reading frames with *Bam*HI, *Bgl*II, *Pst*I, *Nco*I and *Hind*III cloning site, an N-terminal peptide for ProBond purification and blue/white recombinant screening of plaques; Invitrogen (220) can be used.

Mammalian expression vectors contemplated for use in the invention include vectors with inducible promoters, such as the dihydrofolate reductase promoters, any expression vector with a DHFR expression cassette or a DHFR/methotrexate co-amplification vector such as pED (*Pst*I, *Sal*I, *Sba*I, *Sma*I and *Eco*RI cloning sites, with the vector expressing both the cloned gene and DHFR; Kaufman, 1991). Alternatively a glutamine synthetase/methionine sulfoximine co-amplification vector, such as pEE14 (*Hind*III, *Xba*I, *Sma*I, *Sba*I, *Eco*RI and *Bcl*I cloning sites in which the vector expresses glutamine synthetase and the cloned gene; Celltech). A vector that directs episomal expression under the control of the Epstein Barr Virus (EBV) or nuclear antigen (EBNA) can be used such as pREP4 (*Bam*HI, *Sfi*I, *Xho*I, *Not*I, *Nhe*I, *Hind*III, *Nhe*I, *Pvu*II and *Kpn*I cloning sites, constitutive RSV-LTR promoter, hygromycin selectable marker; Invitrogen) pCEP4 (*Bam*HI, *Sfi*I, *Xho*I, *Not*I, *Nhe*I, *Hind*III, *Nhe*I, *Pvu*II and *Kpn*I cloning sites, constitutive hCMV immediate early gene promoter, hygromycin selectable marker; Invitrogen), pMEP4 (*Kpn*I, *Pvu*I, *Nhe*I, *Hind*III, *Not*I, *Xho*I, *Sfi*I, *Bam*HI cloning sites, inducible methallothionein IIa gene promoter, hygromycin selectable marker, Invitrogen), pREP8 (*Bam*HI, *Xho*I, *Not*I, *Hind*III, *Nhe*I and *Kpn*I cloning sites, RSV-LTR promoter, histidinol selectable marker; Invitrogen), pREP9 (*Kpn*I, *Nhe*I, *Hind*III, *Not*I, *Xho*I, *Sfi*I, *Bam*HI cloning sites, RSV-LTR promoter, G418 selectable marker; Invitrogen), and pEBVHis (RSV-LTR promoter, hygromycin selectable marker, N-terminal peptide purifiable via ProBond resin and cleaved by enterokinase; Invitrogen).

Selectable mammalian expression vectors for use in the invention include, but are not limited to, pRc/CMV (*HindIII*, *BstXI*, *NotI*, *SbaI* and *ApaI* cloning sites, G418 selection, Invitrogen), pRc/RSV (*HindII*, *SpeI*, *BstXI*, *NotI*, *XbaI* cloning sites, G418 selection, Invitrogen) and the like. Vaccinia virus mammalian expression vectors (see, for example Kaufman 1991 that can be used in the present invention include, but are not limited to, pSC11 (*SmaI* cloning site, TK- and β -gal selection), pMJ601 (*Sall*, *SmaI*, *AflI*, *NarI*, *BspMII*, *BamHI*, *ApaI*, *NheI*, *SacII*, *KpnI* and *HindIII* cloning sites; TK- and β -gal selection), pTKgptF1S (*EcoRI*, *PstI*, *SaII*, *AccI*, *HindII*, *SbaI*, *BamHI* and *HpaI* cloning sites, TK or XPRT selection) and the like.

Yeast expression systems that can also be used in the present include, but are not limited to, the non-fusion pYES2 vector (*XbaI*, *SphI*, *ShoI*, *NotI*, *GstXI*, *EcoRI*, *BstXI*, *BamHI*, *SacI*, *KpnI* and *HindIII* cloning sites, Invitrogen), the fusion pYESHisA, B, C (*XbaI*, *SphI*, *ShoI*, *NotI*, *BstXI*, *EcoRI*, *BamHI*, *SacI*, *KpnI* and *HindIII* cloning sites, N-terminal peptide purified with ProBond resin and cleaved with enterokinase; Invitrogen), pRS vectors and the like.

Consequently, mammalian and typically human cells, as well as bacterial, yeast, fungi, insect, nematode and plant cells an used in the present invention and may be transfected by the nucleic acid or recombinant vector as defined herein.

Examples of suitable cells include, but are not limited to, VERO cells, HELA cells such as ATCC No. CCL2, CHO cell lines such as ATCC No. CCL61, COS cells such as COS-7 cells and ATCC No. CRL 1650 cells, W138, BHK, HepG2, 3T3 such as ATCC No. CRL6361, A549, PC12, K562 cells, 293 cells, Sf9 cells such as ATCC No. CRL1711 and Cv1 cells such as ATCC No. CCL70.

Other suitable cells that can be used in the present invention include, but are not limited to, prokaryotic host cells strains such as *Escherichia coli*, (e.g., strain DH5- α), *Bacillus subtilis*, *Salmonella typhimurium*, or strains of the genera of *Pseudomonas*, *Streptomyces* and *Staphylococcus*.

Further suitable cells that can be used in the present invention include yeast cells such as those of *Saccharomyces* such as *Saccharomyces cerevisiae*.

Besides the specific isolated complexes, as described above, the present invention relates to and also encompasses SID® polynucleotides. As explained above, for each bait polypeptide, several prey polypeptides may be identified by comparing and selecting the intersection of every isolated fragment that are included in the same polypeptide, as set forth, for example, in described by Szabo et al, *supra*.

The present invention is not limited to the SID® sequences as described in the above paragraph, but also includes fragments of these sequences having at least 12 consecutive nucleic acids, between 12 and 5,000 consecutive nucleic acids and between 12 and 10,000 consecutive nucleic acids and between 12 and 20,000 consecutive nucleic acids, as well as variants thereof. The fragments or variants of the SID® sequences possess at least the same affinity of binding to its protein or polypeptide counterpart, against which it has been initially selected. Moreover this variant and/or fragments of the SID® sequences alternatively can have between 95% and 99.999% sequence identity to its protein or polypeptide counterpart.

According to the present invention the variants can be created by known mutagenesis techniques either *in vitro* or *in vivo*. Such a variant can be created such that it has altered binding characteristics with respect to the target protein and more specifically that the variant binds the target sequence with either higher or lower affinity.

Polynucleotides that are complementary to the above sequences which include the polynucleotides of the SID®'s, their fragments, variants and those that have specific sequence identity are also included in the present invention.

The polynucleotide encoding the SID® polypeptide, fragment or variant thereof can also be inserted into recombinant vectors which are described in detail above.

The present invention also relates to a composition comprising the above-mentioned recombinant vectors containing the SID® polypeptides, fragments or variants thereof, as well as recombinant host cells transformed by the vectors. The recombinant host cells that can be used in the present invention were discussed in greater detail above.

The compositions comprising the recombinant vectors can contain physiological acceptable carriers such as diluents, adjuvants, excipients and any vehicle in which this composition can be delivered therapeutically and can include, but is are not limited to sterile liquids such as water and oils.

In yet another embodiment, the present invention relates to a method of selecting modulating compounds, as well as the modulating molecules or compounds themselves which may be used in a pharmaceutical composition. These modulating compounds may act as a cofactor, as an inhibitor, as antibodies, as tags, as a competitive inhibitor, as an activator or alternatively have agonistic or antagonistic activity on the protein-protein interactions.

The activity of the modulating compound does not necessarily, for example, have to be 100% activation or inhibition. Indeed, even partial activation or inhibition can be achieved that is of pharmaceutical interest.

The modulating compound can be selected according to a method which comprises:

(a) cultivating a recombinant host cell with a modulating compound on a selective medium and a reporter gene the expression of which is toxic for said recombinant host cell wherein said recombinant host cell is transformed with two vectors:

- (i) wherein said first vector comprises a polynucleotide encoding a first hybrid polypeptide having a DNA binding domain;
- (ii) wherein said second vector comprises a polynucleotide encoding a second hybrid polypeptide having a transcriptional activating domain that activates said toxic reporter gene when the first and second hybrid polypeptides interact;

(b) selecting said modulating compound which inhibits or permits the growth of said recombinant host cell.

Thus, the present invention relates to a modulating compound that inhibits the protein-protein interactions of a complex of two polypeptides of columns 1 and 3 of Table 2. The present invention also relates to a modulating compound that activates the protein-protein interactions of a complex of two polypeptides of columns 1 and 3 of Table 2.

In yet another embodiment, the present invention relates to a method of selecting a modulating compound, which modulating compound inhibits the interactions of two polypeptides of columns 1 and 3 of Table 2. This method comprises:

(a) cultivating a recombinant host cell with a modulating compound on a selective medium and a reporter gene the expression of which is toxic for said recombinant host cell wherein said recombinant host cell is transformed with two vectors:

- (i) wherein said first vector comprises a polynucleotide encoding a first hybrid polypeptide having a first domain of an enzyme;
- (ii) wherein said second vector comprises a polynucleotide encoding a second hybrid polypeptide having an enzymatic transcriptional activating domain that activates said toxic reporter gene when the first and second hybrid polypeptides interact;

(b) selecting said modulating compound which inhibits or permits the growth of said recombinant host cell.

In the two methods described above any toxic reporter gene can be utilized including those reporter genes that can be used for negative selection including the URA3 gene, the CYH1 gene, the CYH2 gene and the like.

In yet another embodiment, the present invention provides a kit for screening a modulating compound. This kit comprises a recombinant host cell which comprises a reporter gene the expression of which is toxic for the recombinant host cell. The host cell is transformed with two vectors. The first vector comprises a polynucleotide encoding a first hybrid polypeptide having a DNA binding domain; and a second vector comprises a polynucleotide encoding a second hybrid polypeptide having a transcriptional activating domain that activates said toxic reporter gene when the first and second hybrid polypeptides interact.

In yet another embodiment a kit is provided for screening a modulating compound by providing a recombinant host cell, as described in the paragraph above, but instead of a DNA binding domain, the first vector comprises a first hybrid polypeptide containing a first domain of a protein. The second vector comprises a second polypeptide containing a second part of a complementary domain of a protein that activates the toxic reporter gene when the first and second hybrid polypeptides interact.

In the selection methods described above, the activating domain can be p42 Gal 4, YP16 (HSV) and the DNA-binding domain can be derived from Gal4 or Lex A. The protein or enzyme can be adenylate cyclase, guanylate cyclase, DHFR and the like.

In yet another embodiment, the present invention relates to a pharmaceutical composition comprising the modulating compounds for preventing or treating obesity or metabolic diseases in a human or animal, most preferably in a mammal.

This pharmaceutical composition comprises a pharmaceutically acceptable amount of the modulating compound. The pharmaceutically acceptable amount can be estimated from cell culture assays. For example, a dose can be formulated in animal models to achieve a circulating concentration range that includes or encompasses a concentration point or range having the desired effect in an *in vitro* system. This information can thus be used to accurately determine the doses in other mammals, including humans and animals.

The therapeutically effective dose refers to that amount of the compound that results in amelioration of symptoms in a patient. Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or in experimental animals. For example, the LD50 (the dose lethal to 50% of the population) as well as the ED50 (the dose therapeutically effective in 50% of the population) can be determined using methods known in the art. The dose ratio between toxic and therapeutic effects is the therapeutic index which can be expressed as the ratio between LD 50 and ED50 compounds that exhibit high therapeutic indexes.

The data obtained from the cell culture and animal studies can be used in formulating a range of dosage of such compounds which lies preferably within a range of circulating concentrations that include the ED50 with little or no toxicity.

The pharmaceutical composition can be administered via any route such as locally, orally, systemically, intravenously, intramuscularly, mucosally, using a patch and can be encapsulated in liposomes, microparticles, microcapsules, and the like. The pharmaceutical composition can be embedded in liposomes or even encapsulated.

Any pharmaceutically acceptable carrier or adjuvant can be used in the pharmaceutical composition. The modulating compound will be preferably in a soluble form combined with a pharmaceutically acceptable carrier. The techniques for formulating and administering these compounds can be found in "*Remington's Pharmaceutical Sciences*" Mack Publication Co., Easton, PA, latest edition.

The mode of administration optimum dosages and galenic forms can be determined by the criteria known in the art taken into account the seriousness of the general condition of the mammal, the tolerance of the treatment and the side effects.

The present invention also relates to a method of treating or preventing obesity or metabolic diseases in a human or mammal in need of such treatment. This method comprises administering to a mammal in need of such treatment a pharmaceutically effective amount of a modulating compound which binds to a targeted mammalian or human or adipocyte protein. In a preferred embodiment, the modulating compound is a polynucleotide which may be placed under the control of a regulatory sequence which is functional in the mammal or human.

In yet another embodiment, the present invention relates to a pharmaceutical composition comprising a SID® polypeptide, a fragment or variant thereof. The SID® polypeptide, fragment or variant thereof can be used in a pharmaceutical composition provided that it is endowed with highly specific binding properties to a bait polypeptide of interest.

The original properties of the SID® polypeptide or variants thereof interfere with the naturally occurring interaction between a first protein and a second protein within the cells of the organism. Thus, the SID® polypeptide binds specifically to either the first polypeptide or the second polypeptide.

Therefore, the SID® polypeptides of the present invention or variants thereof interfere with protein-protein interactions between mammalian or human or adipocyte proteins.

Thus, the present invention relates to a pharmaceutical composition comprising a pharmaceutically acceptable amount of a SID® polypeptide or variant thereof, provided

that the variant has the above-mentioned two characteristics; i.e., that it is endowed with highly specific binding properties to a bait polypeptide of interest and is devoid of biological activity of the naturally occurring protein.

In yet another embodiment, the present invention relates to a pharmaceutical composition comprising a pharmaceutically effective amount of a polynucleotide encoding a SID® polypeptide or a variant thereof wherein the polynucleotide is placed under the control of an appropriate regulatory sequence. Appropriate regulatory sequences that are used are polynucleotide sequences derived from promoter elements and the like.

Besides the SID® polypeptides and polynucleotides, the pharmaceutical composition of the present invention can also include a recombinant expression vector comprising the polynucleotide encoding the SID® polypeptide, fragment or variant thereof.

The above described pharmaceutical compositions can be administered by any route such as orally, systemically, intravenously, intramuscularly, intradermally, mucosally, encapsulated, using a patch and the like. Any pharmaceutically acceptable carrier or adjuvant can be used in this pharmaceutical composition.

The SID® polypeptides as active ingredients will be preferably in a soluble form combined with a pharmaceutically acceptable carrier. The techniques for formulating and administering these compounds can be found in "*Remington's Pharmaceutical Sciences*" *supra*.

The amount of pharmaceutically acceptable SID® polypeptides can be determined as described above for the modulating compounds using cell culture and animal models.

Such compounds can be used in a pharmaceutical composition to treat or prevent obesity or any metabolic diseases.

Thus, the present invention also relates to a method of preventing or treating obesity or any metabolic diseases in a mammal said method comprising the steps of administering to a mammal in need of such treatment a pharmaceutically effective amount of:

(1) a SID® polypeptide or a variant thereof which binds to a targeted mammalian or typically human protein; or

(2) or SID® polynucleotide encoding a SID® polypeptide or a variant or a fragment thereof wherein said polynucleotide is placed under the control of a regulatory sequence which is functional in said mammal; or

(3) a recombinant expression vector comprising a polynucleotide encoding a SID® polypeptide which binds to a mammalian or human or adipocyte protein.

In another embodiment the present invention nucleic acids comprising a sequence which encodes the protein and/or functional derivatives thereof are administered to modulate the complex function by way of gene therapy. Any of the methodologies relating to gene therapy available within the art may be used in the practice of the present invention such as those described by Goldspiel et al *Clin. Pharm.* **12** pgs. 488-505 (1993).

Delivery of the therapeutic nucleic acid into a patient may be direct *in vivo* gene therapy (i.e., the patient is directly exposed to the nucleic acid or nucleic acid-containing vector) or indirect *ex vivo* gene therapy (i.e., cells are first transformed with the nucleic acid *in vitro* and then transplanted into the patient).

For example for *in vivo* gene therapy, an expression vector containing the nucleic acid is administered in such a manner that it becomes intracellular; i.e., by infection using a defective or attenuated retroviral or other viral vectors as described, for example in U.S. Patent 4,980,286 or by Robbins et al, *Pharmacol. Ther.* , **80** No. 1 pgs. 35-47 (1998).

The various retroviral vectors that are known in the art are such as those described in Miller et al, *Meth. Enzymol.* **217** pgs. 581-599 (1993) which have been modified to delete those retroviral sequences which are not required for packaging of the viral genome and subsequent integration into host cell DNA. Also adenoviral vectors can be used which are advantageous due to their ability to infect non-dividing cells and such high-capacity adenoviral vectors are described in Kochanek, *Human Gene Therapy*, **10**, pgs. 2451-2459 (1999). Chimeric viral vectors that can be used are those described by Reynolds et al, *Molecular Medicine Today*, pgs. 25 -31 (1999). Hybrid vectors can also be used and are described by Jacoby et al, *Gene Therapy*, **4**, pgs. 1282-1283 (1997).

Direct injection of naked DNA or through the use of microparticle bombardment (e.g., Gene Gun®; Biolistic, Dupont). or by coating it with lipids can also be used in gene therapy. Cell-surface receptors/transfecting agents or through encapsulation in liposomes, microparticles or microcapsules or by administering the nucleic acid in linkage to a peptide which is known to enter the nucleus or by administering it in linkage to a ligand predisposed to receptor-mediated endocytosis (See, Wu & Wu, J. Biol. Chem., 262 pgs. 4429-4432 (1987)) can be used to target cell types which specifically express the receptors of interest.

In another embodiment a nucleic acid ligand compound may be produced in which the ligand comprises a fusogenic viral peptide designed so as to disrupt endosomes, thus allowing the nucleic acid to avoid subsequent lysosomal degradation. The nucleic acid may be targeted *in vivo* for cell specific endocytosis and expression by targeting a specific receptor such as that described in WO92/06180, WO93/14188 and WO 93/20221. Alternatively the nucleic acid may be introduced intracellularly and incorporated within the host cell genome for expression by homologous recombination. See, Zijlstra et al, *Nature*, **342**, pgs. 435-428 (1989).

In *ex vivo* gene a gene is transferred into cells *in vitro* using tissue culture and the cells are delivered to the patient by various methods such as injecting subcutaneously, application of the cells into a skin graft and the intravenous injection of recombinant blood cells such as hematopoietic stem or progenitor cells.

Cells into which a nucleic acid can be introduced for the purposes of gene therapy include, for example, epithelial cells, endothelial cells, keratinocytes, fibroblasts, muscle cells, hepatocytes and blood cells. The blood cells that can be used include, for example, T-lymphocytes, B-lymphocytes, monocytes, macrophages, neutrophils, eosinophils, megakaryocytes, granulocytes, hematopoietic cells or progenitor cells and the like.

In yet another embodiment the present invention relates to protein chips or protein microarrays. It is well known in the art that microarrays can contain more than 10,000 spots of a protein that can be robotically deposited on a surface of a glass slide or nylon filter. The proteins attach covalently to the slide surface, yet retain their ability to interact with other proteins or small molecules in solution. In some instances the protein samples can be made to adhere to glass slides by coating the slides with an aldehyde-containing

reagent that attaches to primary amines. A process for creating microarrays is described, for example by MacBeath and Schreiber in *Science*, Volume 289, Number 5485, pgs, 1760-1763 (2000) or Service, *Science*, Vol, 289, Number 5485 pg. 1673 (2000). An apparatus for controlling, dispensing and measuring small quantities of fluid is described, for example, in U.S. Patent No. 6,112,605.

The present invention also provides a record of protein-protein interactions, PIM®'s, SID®'s and any data encompassed in the following Tables. It will be appreciated that this record can be provided in paper or electronic or digital form.

In order to fully illustrate the present invention and advantages thereof, the following specific examples are given, it being understood that the same are intended only as illustrative and in nowise limitative.

EXAMPLES

EXAMPLE 1: Preparation of a collection of random-primed cDNA fragments

1.A. Collection preparation and transformation in *Escherichia coli*

1.A.1. Random-primed cDNA fragment preparation

For each mRNA sample (human placenta, undifferentiated PAZ6 adipocytes or differentiated PAZ6 adipocytes), random-primed cDNA was prepared from 5 µg of polyA+ mRNA using a TimeSaver cDNA Synthesis Kit (Amersham Pharmacia Biotech) and with 5 µg of random N9-mers according to the manufacturer's instructions. Following phenolic extraction, the cDNA was precipitated and resuspended in water. The resuspended cDNA was phosphorylated by incubating in the presence of T4 DNA Kinase (Biolabs) and ATP for 30 minutes at 37°C. The resulting phosphorylated cDNA was then purified over a separation column (Chromaspin TE 400, Clontech), according to the manufacturer's protocol.

1.A.2. Ligation of linkers to blunt-ended cDNA

Oligonucleotide HGX931 (5' end phosphorylated) 1 µg/µl and HGX932 1µg/µl.

Sequence of the oligo HGX931: 5'-GGGCCACGAA-3' (SEQ ID No. 61)

Sequence of the oligo HGX932: 5'-TTCGTGGCCCCTG-3' (SEQ ID No. 62)

Linkers were preincubated (5 minutes at 95°C, 10 minutes at 68°C, 15 minutes at 42°C) then cooled down at room temperature and ligated with cDNA fragments at 16°C overnight.

Linkers were removed on a separation column (Chromaspin TE 400, Clontech), according to the manufacturer's protocol.

1.A.3. Vector preparation

Plasmid pP6 (see Figure 10) was prepared by replacing the *SpeI/XhoI* fragment of pGAD3S2X with the double-stranded oligonucleotide:

5'CTAGCCATGGCCGCAGGGGCCGCGGCCGCACTAGTGGGGATCCTTAATTAAAGG
GCCACTGGGGCCCCC
GGTACCGGCGTCCCCGGCGCCGGCGTGATCACCCCTAGGAATTAATTTCCCGGTGA
CCCCGGGGGAGCT 3' (SEQ ID No. 63)

The pP6 vector was successively digested with *SfiI* and *BamHI* restriction enzymes (Biolabs) for 1 hour at 37°C, extracted, precipitated and resuspended in water. Digested plasmid vector backbones were purified on a separation column (Chromaspin TE 400, Clontech), according to the manufacturer's protocol.

1.A.4. Ligation between vector and insert of cDNA

The prepared vector was ligated overnight at 15°C with the blunt-ended cDNA described in section 2 using T4 DNA ligase (Biolabs). The DNA was then precipitated and resuspended in water.

1.A.5. Library transformation in *Escherichia coli*

The DNA from section 1.A.4 was transformed into Electromax DH10B electrocompetent cells (Gibco BRL) with a Cell Porator apparatus (Gibco BRL). 1 ml SOC medium was added and the transformed cells were incubated at 37°C for 1 hour. 9 mls of SOC medium per tube was added and the cells were plated on LB+ampicillin medium. The colonies were scraped with liquid LB medium, aliquoted and frozen at -80°C.

The obtained collections of recombinant cell clones were named: HGXBPLARP1 (placenta), HGXPZURP1 (undifferentiated PAZ6 adipocytes) and HGXPZDRP1 (differentiated PAZ6 adipocytes).

1.B. Collection transformation in *Saccharomyces cerevisiae*

The *Saccharomyces cerevisiae* strain (Y187 (MAT α Gal4 Δ Gal80 Δ ade2-101, his3, leu2-3, -112, trp1-901, ura3-52 URA3::UASGAL1-LacZ Met)) was transformed with the cDNA library.

The plasmid DNA contained in *E. coli* were extracted (Qiagen) from aliquoted *E. coli* frozen cells (1.A.5.). *Saccharomyces cerevisiae* yeast Y187 in YPGlu were grown.

Yeast transformation was performed according to standard protocol (Giest *et al.* Yeast, 11, 355-360, 1995) using yeast carrier DNA (Clontech). This experiment leads to 10⁴ to 5 x 10⁴ cells/ μ g DNA. 2 x 10⁴ cells were spread on DO-Leu medium per plate. The cells were aliquoted into vials containing 1 ml of cells and frozen at -80°C.

The obtained collections of recombinant cell clones are named: HGXYPLARP1 (placenta), HGXYPZURP1 (undifferentiated PAZ6 adipocytes) and HGXYPZDRP1(differentiated PAZ6 adipocytes).

1.C. Construction of bait plasmids

For fusions of the bait protein to the DNA-binding domain of the GAL4 protein of *S. cerevisiae*, bait fragments were cloned into plasmid pB6. For fusions of the bait protein to the DNA-binding domain of the LexA protein of *E. coli*, bait fragments were cloned into plasmid pB20.

Plasmid pB6 (see Figure 3) was prepared by replacing the *Nco*1/*Sal*1 polylinker fragment of pAS $\Delta\Delta$ with the double-stranded DNA fragment:

```

5'
CATGGCCGGACGGGCGCGGCCGCACTAGTGGGGATCCTTAATTAAGGGCCACT
GGGGCCCCC 3'
3'
CGGCCTGCCCCGGCGCCGGCGTGATCACCCCTAGGAATTAATTTCCCGGTGACCCC
GGGGGAGCT 5' (SEQ ID No. 64)

```

Plasmid pB20 (see Figure 6) was prepared by replacing the *Eco*RI/*Pst*I polylinker fragment of pLex10 with the double-stranded DNA fragment:

```

5'
AATTCGGGGCCGGACGGGCGCGGCCGCACTAGTGGGGATCCTTAATTAAGGGCC
ACTGGGGCCCCCTCGACCTGCA 3'

```


3'

CCCCCGGCCTGCCCGGCGCCGGCGTGATCACCCCTAGGAATTAATTCCCGGTGAC
CCCGGGGAGCTGG 5' (SEQ ID No. 65)

The amplification of the bait ORF was obtained by PCR using the Pfu proof-reading *Taq* polymerase (Stratagene), 10 pmol of each specific amplification primer and 200 ng of plasmid DNA as template.

The PCR program was set up as follows :

94°	45"	} x 30 cycles
94°	45"	
48°	45"	
72°	6'	
72°	10'	
15°	∞	

The amplification was checked by agarose gel electrophoresis.

The PCR fragments were purified with Qiaquick column (Qiagen) according to the manufacturer's protocol.

Purified PCR fragments were digested with adequate restriction enzymes.

The PCR fragments were purified with Qiaquick column (Qiagen) according to the manufacturer's protocol.

The digested PCR fragments were ligated into an adequately digested and dephosphorylated bait vector (pB6 or pB20) according to standard protocol (Sambrook *et al.*) and were transformed into competent bacterial cells. The cells were grown, the DNA extracted and the plasmid was sequenced.

Example 2 : Screening the collection with the two-hybrid in yeast system

2.A. The mating protocol

The mating two-hybrid in yeast system (as described by Legrain *et al.*, *Nature Genetics*, vol. 16, 277-282 (1997), *Toward a functional analysis of the yeast genome through exhaustive two-hybrid screens*) was used for its advantages but one could also screen the cDNA collection in classical two-hybrid system as described in Fields *et al.* or in a yeast reverse two-hybrid system.

The mating procedure allows a direct selection on selective plates because the two fusion proteins are already produced in the parental cells. No replica plating is required.

This protocol was written for the use of the library transformed into the Y187 strain.

For bait proteins fused to the DNA-binding domain of GAL4, bait-encoding plasmids were first transformed into *S. cerevisiae* (CG1945 strain (MATa Gal4-542 Gal180-538 ade2-101 his3Δ200, leu2-3,112, trp1-901, ura3-52, lys2-801, URA3::GAL4 17mers (X3)-CyC1TATA-LacZ, LYS2::GAL1UAS-GAL1TATA-HIS3 CYH^R)) according to step 1.B. and spread on DO-Trp medium.

For bait proteins fused to the DNA-binding domain of LexA, bait-encoding plasmids were first transformed into *S. cerevisiae* (L40Δgal4 strain (MATa ade2, trp1-901, leu2 3,112, lys2-801, his3Δ200, LYS2::(*lexAop*)₄-HIS3, ura3-52::URA3 (*lexAop*)₈-LacZ, GAL4::Kan^R)) according to step 1.B. and spread on DO-Trp medium.

Day 1, morning : preculture

The cells carrying the bait plasmid obtained at step 1.C. were precultured in 20 ml DO-Trp medium and grown at 30°C with vigorous agitation.

Day 1, late afternoon : culture

The OD_{600nm} of the DO-Trp pre-culture of cells carrying the bait plasmid pre-culture was measured. The OD_{600nm} must lie between 0.1 and 0.5 in order to correspond to a linear measurement.

50 ml DO-Trp at OD_{600nm} 0.006/ml was inoculated and grown overnight at 30°C with vigorous agitation.

Day 2 : mating

medium and plates

- 1 YPGlu 15cm plate
- 50 ml tube with 13 ml DO-Leu-Trp-His
- 100 ml flask with 5 ml of YPGlu
- 8 DO-Leu-Trp-His plates
- 2 DO-Leu plates
- 2 DO-Trp plates

2 DO-Leu-Trp plates

The OD_{600nm} of the DO-Trp culture was measured. It should be around 1.

For the mating, twice as many bait cells as library cells were used. To get a good mating efficiency, one must collect the cells at 10^8 cells per cm².

The amount of bait culture (in ml) that makes up 50 OD_{600nm} units for the mating with the prey library was estimated.

A vial containing the HGXYCDNA1 library was thawed slowly on ice. 1.0ml of the vial was added to 5 ml YPGlu. Those cells were recovered at 30°C, under gentle agitation for 10 minutes.

Mating

The 50 OD_{600nm} units of bait culture was placed into a 50 ml falcon tube.

The HGXYCDNA1 library culture was added to the bait culture, then centrifuged, the supernatant discarded and resuspended in 1.6ml YPGlu medium.

The cells were distributed onto two 15cm YPGlu plates with glass beads. The cells were spread by shaking the plates. The plate cells-up at 30°C for 4h30min were incubated.

Collection of mated cells

The plates were washed and rinsed with 6ml and 7ml respectively of DO-Leu-Trp-His. Two parallel serial ten-fold dilutions were performed in 500µl DO-Leu-Trp-His up to 1/10,000. 50µl of each 1/10000 dilution was spread onto DO-Leu and DO-trp plates and 50µl of each 1/1000 dilution onto DO-Leu-Trp plates. 22.4ml of collected cells were spread in 400µl aliquots on DO-Leu-Trp-His+Tet plates.

Day 4

Clones that were able to grow on DO-Leu-Trp-His+Tetracyclin were then selected. This medium allows one to isolate diploid clones presenting an interaction.

The His⁺ colonies were counted on control plates.

The number of His⁺ cell clones will define which protocol is to be processed :

Upon $60 \cdot 10^6$ Trp⁺Leu⁺ colonies :

- if the number His⁺ cell clones <285 : then use the process luminometry protocol on all colonies
- if the number of His⁺ cell clones > 285 and <5000: then process via overlay and then luminometry protocols on blue colonies (2.B and 2.C).
- if number of His⁺ cell clones >5000 : repeat screen using DO-Leu-Trp-His+Tetracyclin plates containing 3-aminotriazol.

2.B. The X-Gal overlay assay

The X-Gal overlay assay was performed directly on the selective medium plates after scoring the number of His⁺ colonies.

Materials

A waterbath was set up. The water temperature should be 50°C.

- 0.5 M Na₂HPO₄ pH 7.5.
- 1.2% Bacto-agar.
- 2% X-Gal in DMF.
- Overlay mixture : 0.25 M Na₂HPO₄ pH7.5, 0.5% agar, 0.1% SDS, 7% DMF (LABOSI), 0.04% X-Gal (ICN). For each plate, 10 ml overlay mixture are needed.
- DO-Leu-Trp-His plates.
- Sterile toothpicks.

Experiment

The temperature of the overlay mix should be between 45°C and 50°C. The overlay-mix was poured over the plates in portions of 10 ml. When the top layer was settled, they were collected. The plates were incubated overlay-up at 30°C and the time was noted. Blue colonies were checked for regularly. If no blue colony appeared, overnight incubation was performed. Using a pen the number of positives was marked. The positives colonies were streaked on fresh DO-Leu-Trp-His plates with a sterile toothpick.

2.C. The luminometry assay

His⁺ colonies were grown overnight at 30°C in microtiter plates containing DO-Leu-Trp-His+Tetracyclin medium with shaking. The day after, the overnight culture was diluted 15 times into a new microtiter plate containing the same medium and was incubated for 5 hours at 30°C with shaking. The samples were diluted 5 times and read OD_{600nm}. The samples were diluted again to obtain between 10,000 and 75,000 yeast cells/well in 100 µl final volume.

Per well, 76 µl of One Step Yeast Lysis Buffer (Tropix) was added, 20 µl SapphireII Enhancer (Tropix), 4 µl Galacton Star (Tropix) and incubated 40 minutes at 30°C. The β-Gal read-out (L) was measured using a Luminometer (Trilux, Wallach). The value of (OD_{600nm} × L) was calculated and interacting preys having the highest values were selected.

At this step of the protocol, diploid cell clones presenting interaction were isolated. The next step was now to identify polypeptides involved in the selected interactions.

Example 3 : Identification of positive clones

3.A. PCR on yeast colonies

Introduction

PCR amplification of fragments of plasmid DNA directly on yeast colonies is a quick and efficient procedure to identify sequences cloned into this plasmid. It is directly derived from a published protocol (Wang H. et al., *Analytical Biochemistry*, **237**, 145-146, (1996)). However, it is not a standardized protocol and it varies from strain to strain and it is dependent of experimental conditions (number of cells, *Taq* polymerase source, etc). This protocol should be optimized to specific local conditions.

Materials

- For 1 well, PCR mix composition was :

32.5 μ l water,

5 μ l 10X PCR buffer (Pharmacia),

1 μ l dNTP 10 mM,

0.5 μ l *Taq* polymerase (5u/ μ l) (Pharmacia),

0.5 μ l oligonucleotide ABS1 10 pmole/ μ l: 5'-GCGTTTGGAATCACTACAGG-3', (SEQ ID No.66)

0.5 μ l oligonucleotide ABS2 10 pmole/ μ l: 5'-CACGATGCACGTTGAAGTG-3'. (SEQ ID No. 67)

- 1 N NaOH.

Experiment

The positive colonies were grown overnight at 30°C on a 96 well cell culture cluster (Costar), containing 150 μ l DO-Leu-Trp-His+Tetracyclin with shaking. The culture was resuspended and 100 μ l was transferred immediately on a Thermowell 96 (Costar) and centrifuged for 5 minutes at 4,000 rpm at room temperature. The supernatant was removed. 5 μ l NaOH was added to each well and shaken for 1 minute.

The Thermowell was placed in the thermocycler (GeneAmp 9700, Perkin Elmer) for 5 minutes at 99.9°C and then 10 minutes at 4°C. In each well, the PCR mix was added and shaken well.

The PCR program was set up as followed :

94°C 3 minutes

94°C	30 seconds	
53°C	1 minute 30 seconds	x 35 cycles
72°C	3 minutes	
72°C	5 minutes	
15°C	∞	

The quality, the quantity and the length of the PCR fragment was checked on an agarose gel. The length of the cloned fragment was the estimated length of the PCR fragment minus 300 base pairs that corresponded to the amplified flanking plasmid sequences.

3.B. Plasmids rescue from yeast by electroporation

Introduction

The previous protocol of PCR on yeast cell may not be successful, in such a case, plasmids from yeast by electroporation can be rescued. This experiment allows the recovery of prey plasmids from yeast cells by transformation of *E. coli* with a yeast cellular extract. The prey plasmid can then be amplified and the cloned fragment can be sequenced.

Materials

Plasmid rescue

Glass beads 425-600 μm (Sigma)

Phenol/chloroform (1/1) premixed with isoamyl alcohol (Amresco)

Extraction buffer : 2% Triton X100, 1% SDS, 100 mM NaCl, 10 mM TrisHCl pH 8.0, 1 mM EDTA pH 8.0.

Mix ethanol/ NH_4Ac : 6 volumes ethanol with 7.5 M NH_4 Acetate, 70% Ethanol and yeast cells in patches on plates.

Electroporation

SOC medium

M9 medium

Selective plates : M9-Leu+Ampicillin

2 mm electroporation cuvettes (Eurogentech)

Experiment

Plasmid rescue

The cell patch on DO-Leu-Trp-His was prepared with the cell culture of section 2.C. The cell of each patch was scraped into an Eppendorf tube, 300 μl of glass beads

was added in each tube, then, 200 μ l extraction buffer and 200 μ l phenol:chloroform:isoamyl alcohol (25:24:1) was added.

The tubes were centrifuged for 10 minutes at 15,000 rpm.

180 μ l supernatant was transferred to a sterile Eppendorf tube and 500 μ l each of ethanol/ NH_4Ac was added and the tubes were vortexed. The tubes were centrifuged for 15 minutes at 15,000 rpm at 4°C. The pellet was washed with 200 μ l 70% ethanol and the ethanol was removed and the pellet was dried. The pellet was resuspended in 10 μ l water. Extracts were stored at -20°C.

Electroporation

Materials : Electrocompetent MC1066 cells prepared according to standard protocols (Sambrook et al. *supra*).

1 μ l of yeast plasmid DNA-extract was added to a pre-chilled Eppendorf tube, and kept on ice.

1 μ l plasmid yeast DNA-extract sample was mixed and 20 μ l electrocompetent cells was added and transferred in a cold electroporation cuvette.

Set the Biorad electroporator on 200 ohms resistance, 25 μ F capacity; 2.5 kV. Place the cuvette in the cuvette holder and electroporate.

1 ml of SOC was added into the cuvette and the cell-mix was transferred into a sterile Eppendorf tube. The cells were recovered for 30 minutes at 37°C, then spun down for 1 minute at 4,000 x g and the supernatant was poured off. About 100 μ l medium was kept and used to resuspend the cells and spread them on selective plates (e.g., M9-Leu plates). The plates were then incubated for 36 hours at 37°C.

One colony was grown and the plasmids were extracted. Check for the presence and size of the insert through enzymatic digestion and agarose gel electrophoresis. The insert was then sequenced.

Example 4 : Protein-protein interaction

For each bait, the previous protocol leads to the identification of prey polynucleotide sequences. Using a suitable software program (e.g., Blastwun, available on the Internet site of the University of Washington : <http://bioweb.pasteur.fr/seqanal/interfaces/blastwu.html>) the identity of the mRNA transcript that is encoded by the prey fragment may be determined and whether the

fusion protein encoded is in the same open reading frame of translation as the predicted protein or not.

Alternatively, prey nucleotide sequences can be compared with one another and those which share identity over a significant region (60nt) can be grouped together to form a contiguous sequence (Contig) whose identity can be ascertained in the same manner as for individual prey fragments described above.

Example 5 : Identification of SID®

By comparing and selecting the intersection of all isolated fragments that are included in the same polypeptide, one can define the Selected Interacting Domain (SID®) is determined as illustrated in Figure 15.

Example 6: Making of polyclonal and monoclonal antibodies

The protein-protein complex of columns 1 and 3 of Table 2 was injected into mice and polyclonal and monoclonal antibodies were made following the procedure set forth in Sambrook et al *supra*.

More specifically, mice are immunized with an immunogen comprising the above mentioned complexes conjugated to keyhole limpet hemocyanin using glutaraldehyde or EDC as is well known in the art. The complexes can also be stabilized by crosslinking as described in WO 00/37483. The immunogen is then mixed with an adjuvant. Each mouse receives four injections of 10 ug to 100 ug of immunogen, and after the fourth injection, blood samples are taken from the mice to determine if the serum contains antibodies to the immunogen. Serum titer is determined by ELISA or RIA. Mice with sera indicating the presence of antibody to the immunogen are selected for hybridoma production.

Spleens are removed from immune mice and single-cell suspension is prepared (Harlow et al 1988). Cell fusions are performed essentially as described by Kohler et al.. Briefly, P365.3 myeloma cells (ATTC Rockville, Md) or NS-1 myeloma cells are fused with spleen cells using polyethylene glycol as described by Harlow et al (1989). Cells are plated at a density of 2×10^5 cells/well in 96-well tissue culture plates. Individual wells are examined for growth and the supernatants of wells with growth are tested for the presence of complex-specific antibodies by ELISA or RIA using the protein-protein complex of columns 1 and 3 of Table 2 as a target protein. Cells in positive wells are expanded and subcloned to establish and confirm monoclonality.

Clones with the desired specificities are expanded and grown as ascites in mice or in a hollow fiber system to produce sufficient quantities of antibodies for characterization and assay development. Antibodies are tested for binding to bait polypeptide of column 1 of Table 2 alone or to prey polypeptide of column 3 of Table 2 alone, to determine which are specific for the protein-protein complex of columns 1 and 3 of Table 2 as opposed to those that bind to the individual proteins.

Monoclonal antibodies against each of the complexes set forth in columns 1 and 3 of Table 2 are prepared in a similar manner by mixing specified proteins together, immunizing an animal, fusing spleen cells with myeloma cells and isolating clones which produce antibodies specific for the protein complex, but not for individual proteins.

Example 7: Modulating compounds identification

Each specific protein-protein complex of columns 1 and 3 of Table 2 may be used to screen for modulating compounds.

One appropriate construction for this modulating compound screening may be:

- bait polynucleotide inserted in pB6;
- prey polynucleotide inserted in pP6;
- transformation of these two vectors in a permeable yeast cell;
- growth of the transformed yeast cell on a medium containing compound to be tested,
- and observation of the growth of the yeast cells.

The following results obtained from these Examples, as well as the teachings in the specification are set forth in the Tables below.

Example 8

Materials and Methods

1. Plasmid constructions, transfections and cell culture.

The GW1-HA-MUPP1 plasmid containing the coding region of MUPP1 (multi-PDZ - domain protein) has been obtained by Dr. Javier (Barritt et al. J Cell Biochem 79:213-224 (2000) and Lee et al. J Virol 74: 9680-9693 (2000). MTR-YFP and MTR-Rluc fusion proteins were constructed by ligating the YFP and the Rluc moieties at the C-terminal end of the receptors. For this, the coding regions of MT1R and MT2R were inserted into the cloning sites of the pRL-CMV vector (Promega, Madison, WI) in phase with the *Renilla* luciferase gene or cloned in phase with the YFP coding region of the Cytogem®-Topaze (pGFPTpz-N1) vector (Packard, Meriden, CT). Stop codons were then deleted by site-directed mutagenesis. All constructs were verified by sequencing.

HEK 293 cells were grown in DMEM supplemented with 10% (v/v) FBS, 4.5 g/liter glucose, 100 U/ml penicillin, 0.1 mg/ml streptomycin, 1 mM glutamine (all from Life Technologies (Gaithersburg, MD)). Transient and stable transfections were performed using the transfection reagent FuGene 6 (Roche, Basel, Switzerland) according to supplier instructions.

2. Membrane preparation, solubilization and immunoprecipitation.

Cells were put on ice, washed once with ice-cold PBS and lysed in 350 µl of lysis buffer (25 mM Hepes, 150mM NaCl, 2 mM EDTA, 15 mM β-glycerophosphate, 2 mM Na₃VO₄, 10 mM NaF, 5 µg/ml leupeptin, 10 µg/ml pepstatin, 10 µg/ml benzamidin, 1 mM AEBSF) containing 1 % digitonin for 4 h. The volume was adjusted to 1 ml with lysis buffer without digitonin, and the lysate centrifuged at 18,000 x g for 30 min at 4°C. The supernatant (850 µl) was added to 3 µg of the Flag-specific M2 antibody (Sigma, St Louis, MO) pre-adsorbed on Protein G. After 16 h incubation, immunoadsorbed material was pelleted by centrifugation and washed three times with 1 ml lysis buffer without detergent.

3. SDS-PAGE / Immunoblotting

Whole cell lysates or immunoprecipitates were denatured in 62.5 mM Tris/HCl (pH 6.8), 5% SDS, 10% glycerol, 0.05% bromophenol blue at room temperature. Proteins

were separated by 7 % SDS-PAGE and transferred to nitrocellulose. Immunoblot analysis was carried out with the polyclonal anti-MUPP1 (Barritt et al. J Cell Biochem 79:213-224 (2000) and Lee et al. J Virol 74: 9680-9693 (2000). Immunoreactivity was revealed using a goat anti-rabbit secondary antibody coupled to horseradish peroxidase and the ECL chemi-luminescent reagent (Amersham, Aylesbury, UK).

4. Radioligand Binding Experiments

Whole cell radioligand binding assays were performed as described (Brydon, L., Rocka, F., Petit, L., de Coppet, P., Tissot, M., Barrett, P., Morgan, P. J., Nanoff, C., Strosberg, A. D., and Jockers, R. (1999) *Mol Endocrinol* **13**, 2025- 2038). 2(¹²⁵I)-iodomelatonin (¹²⁵I-Mel) was used at 400 pM for MTR (NEN, Boston, MA). Specific binding was defined as binding displaced by 10 μ M melatonin (MTR) (Sigma, St Louis, MO).

5. BRET Assay.

The interaction between melatonin receptors and MUPP1 has been evaluated by a protein-protein interaction assay based on the bioluminescence resonance energy transfer (BRET) technology described by Xu et al. (Xu, Y., Piston, D. W., and Johnson, C. H. (1999) *Proc Natl Acad Sci U S A* **96**, 151-156). Cells were transfected with constant amounts of fusion receptors (MT1R-Rluc/MT1R-YFP, MT2R-Rluc/MT2R-YFP or MT1R-Rluc/MT2R-YFP) at a 1:1 ratio (0.4 μ g of each DNA) and 0.4 μ g or increasing amounts of GW1-HA-MUPP1 plasmid. Forty-eight hours post-transfection, HEK 293 cells were detached and washed with PBS. 1.2×10^5 cells were distributed in a 96-well microplate at 25°C. Coelenterazine h (Molecular Probes, Eugene, OR) was added at a final concentration of 5 μ M and readings were performed with a lumino/fluorometer (Fusion™, Packard Instrument Company, Meriden, CT) that allows the sequential integration of luminescence signals detected with two filter settings (Rluc filter : 485 ± 10 nm; YFP filter : 530 ± 12.5 nm). The BRET ratio was defined as the difference of the emission at 530 nm/485 nm) of co-transfected Rluc and YFP fusion proteins and the emission at 530 nm/485 nm of the Rluc fusion protein alone. Results were expressed in milliBRET Units (mBU), 1 mBRET Unit corresponding to the BRET ratio values multiplied by 1000. The amount of Rluc and YFP expressed was determined for each condition. Maximal luciferase activity was used to determine the amount of Rluc fusion receptors and the

fluorescence obtained upon exogenous YFP excitation to determine the amount of YFP fusion receptors.

Results

The interaction between MT1R and MUPP1 has been confirmed by co-immunoprecipitation experiments in HEK 293 cells transfected with the MT1R cDNA in the presence or absence of MUPP1 cDNA. MUPP1 expression was verified in whole cell lysates by Western blotting. Flag-tagged MT1 receptors were immunoprecipitated with anti Flag antibodies and MUPP1 was detected in precipitates with an anti-MUPP1 antibody on Western blots (Fig. 17).

The interaction between MT1R and MUPP1 was also verified in BRET experiments. Expression of MUPP1 decreased the energy transfer between MT1R-Rluc and MT1R-YFP in a dose dependent manner (Fig. 18 B). The energy transfer between MT1R-Rluc and MT2R-YFP was also inhibited although to a lesser extent (Fig. 18 A). The decrease of the energy transfer may be explained by the specific interaction of MUPP1 with the carboxy-terminus of the MT1R. This interaction changes the position of the luciferase and YFP molecule, which are fused to the carboxy-terminal tail of the receptors, and thus decreases the energy transfer. The transfer between MT2R-Rluc and MT2R-YFP was insensitive to MUPP1 expression confirming the specificity of the interaction (Fig. 18 A).

Example 9

β arrestin2/Oct-1

Studies were carried out to investigate the potential functionality of the interaction between β arrestin2 and Oct-1, identified by the yeast two-hybrid system. Oct-1 is a ubiquitously expressed member of the POU (Pit-1, Oct-1, unc-86) family of transcription factors and is involved in the regulation of a wide variety of genes implicated in cell cycle regulation, development and hormonal signals. It has been demonstrated that Oct-1 can act both as a transcriptional activator and inhibitor for certain genes. Oct-1 has a nuclear localization within the cell, whereas β arrestin2 is cytoplasmic. Recently, however, it was demonstrated that β arrestin2 shuttles between the cytoplasm and the nucleus in studies using Leptomycin B (an inhibitor of nuclear export; Scott *et al.*, manuscript in preparation). The molecular determinants underlying this nucleocytoplasmic shuttling phenotype and mapped a nuclear export signal (NES) in β arrestin2 was therefore characterized.

A reporter gene strategy was used to determine if the expression of wild-type β arrestin2 or a point mutant of β arrestin2 rendering the NES inactive (β arrestin2 Δ NES) and allowing nuclear accumulation of β arrestin2, would have any effect on Oct-1-driven gene expression. Cos-7 cells (which express low levels of endogenous Oct-1) were transfected with a luciferase reporter gene under the control of 8 copies of the octamer binding motif, the binding motif for Oct-1 (8 x Oct-Luc, a kind gift from P. Matthias, Friedrich Miescher-Institut, Basel, Switzerland). The cells were also transfected with β arrestin2 or β arrestin2 Δ NES alone or in combination with Oct-1.

The results (shown in Figure 19) indicate that β arrestin2 has an inhibitory affect on Oct-1-mediated gene expression. Removal of the NES in β arrestin2, however doesn't r to alter this inhibition.

While the invention has been described in terms of the various preferred embodiments, the skilled artisan will appreciate that various modifications, substitutions, omissions and changes may be made without departing from the scope thereof. Accordingly, it is intended that the present invention be limited by the scope of the following claims, including equivalents thereof.

Table 1 : bait function and sequence

1-Bait Protein Name	2-Bait Sequences		3-GenBank Access Number	4-Chromosome Number	5-Start Nucleotide	6-Stop Nucleotide
	2.1-Bait Nucleic Acid Sequence	2.2-Bait Aminoacid sequence				
Human Skp1 : Part of SCF (Skp1/Cullin/F-box) complexes which act as E3 Ubiquitin ligases.	SEQ ID 1 (see sequence in GenBank)	SEQ ID 2 (see sequence in GenBank)	995823	7	94	585
Human Splicing Factor 1	SEQ ID 3 (see sequence in GenBank)	SEQ ID 4 (see sequence in GenBank)	1620402	11	295	2211
mouse p53 : Tumour suppressor protein	SEQ ID 5 (see sequence in GenBank)	SEQ ID 6 (see sequence in GenBank)	2961246		26	1198
Human beta-TCP1 : F-box containing protein with 7 WD40 repeats; Part of SCF (Skp1/Cullin/F-box) complex E3 ubiquitin ligase; Implicated in the degradation of b-catenin and IkBa	SEQ ID 7 (see sequence in GenBank)	SEQ ID 8 (see sequence in GenBank)	4502476	10	70	1779
human Rac1 : Member of Ras subfamily of RAS small GTPases	SEQ ID 9 (see sequence in GenBank)	SEQ ID 10 (see sequence in GenBank)	9845510	7	1	579
Human uracil DNA glycosylase : Uracil DNA glycosylase implicated in DNA repair	SEQ ID 11 (see sequence in GenBank)	SEQ ID 12 (see sequence in GenBank)	6224978	12	107	1021

Human b2 adrenergic receptor : Oncogene	SEQ ID 13 : TCTTTCAGGAGGCCAAAGGCA GCTCCAGAAGATTGACAAATCT GAGGGCCGCTTCCATGTCCAG AACCTTAGCCAGGTGGAGCAG GATGGCGGACGGGGCATGGA CTCCGCAGATCTTCCAAGTTCT GCTTGAAGGAGCACAAAGCCC TCAAGTGA	SEQ ID 14 : VFQEAQRQLQKIDKSEGRFHVQN LSQVEQDGRGTGHLRRSSKFCLK EHKALK	#	#	#	
Human b2 adrenergic receptor : Oncogene	SEQ ID 15 : CGGAGCCCAGATTTTCAGGATTG CCTTCCAGGAGCTTCTGTGCCT GCGCAGGTCTTCTTTGAAGGCC TATGGCAATGGCTACTCCAGCA ACGGCAACACAGGGGAGCAGA GTGGATATCACGTGGAAACAGG AGAAAGAAAAATAAACTGCTGTG TGAAGACCTCCAGGCCACCGGA AGACTTTGTGGGCCATCAAGGT ACTGTGCCTAGCGATAACATTG ATTCACAAGGGAGGAATTGTAG TACAAATGACTCACTGCTATAA	SEQ ID 16 : RSPDFRIAFQELLCLRRSSLKAYG NGYSSNGNTGEQSGYHVEQEKE NKLLCEDLPGTEDFVGHQGTGPS DNIDSQGRNCSTNDSLL	#	#	#	

Human b2 adrenergic receptor : Oncogene	SEQ ID 17 : GTCTTTCAGGAGGCCAAAAGG CAGCTCCAGAAAGATTGACAAAT CTGAGGGCCGCTTCCATGTCC AGAACCCTAGCCAGGTGGAGC AGGATGGGCGGACGGGGCATG GACTCCGCAGATCTTCCAAGTT CTGCTTGAAGGAGCACAAAAGC CCTCAAGGGATCCCGGAGCCC AGATTTCAGGATTGCCCTCCAG GAGCTTCTGTGCCTGCGCAGG TCTTCTTTGAAGGCCTATGGCA ATGGCTACTCCAGCAACGGCAA CACAGGGGAGCAGAGTGGATA TCACGTGGAAACAGGAGAAAGA AAATAAACTGCTGTGTGAAGAC CTCCAGGCACGGAAGACTTT GTGGGCCATCAAGGTACTGTG CCTAGCGATAACATTGATTCAC AAGGGAGGAATTGTAGTACAAA TGACTCACTGCTATAA	SEQ ID 18 : VFQEAQRQLQKIDKSEGRFHVQN LSQVEQDGRGTGHLRRSSKFCLK EHKALKGSRSPDFRIAFQELLCLR RSSLKAYGNGYSSNGNTGEQSG YHVEQEKENKLLCEDLPGTEDFV GHQGTVPDNDIDSQGRNCSTNDS LL	#	#	#	#
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16638010_072302

Human b2 adrenergic receptor : Oncogene	SEQ ID 19 : GTCCTTCAGGAGGCCAAAAGG CAGCTCCAGAAAGATTGACAAAT CTGAGGGCCGCTTCCATGTCC AGAACCTTAGCCAGGTGGAGC AGGATGGCGGACGGGGCATG GACTCCGAGATCTTCCAAGTT CTGCTTGAAGGAGCACAAAGC CCTCAAGACGTTAGGCATCATC ATGGGCACCTTCACCCCTCTGCT GGCTGCCCTTCTTCATCGTTAA CATTGTGCATGTGATCCAGGAT AACCTCATCCGTAAGGAAGTTT ACATCCTCCTAAATTGGATAGG CTATGTCAATTCTGGTTTCAATC CCCTTATCTACTGCCGGAGCCC AGATTTCAGGATTGCCCTTCCAG GAGCTTCTGTGCCTGCGCAGG TCCTCTTTGAAGGCCTATGGCA ATGGCTACTCCAGCAACGGCAA CACAGGGGAGCAGAGTGGATA TCACGTGGAACAGGAGAAAGA AAATAAACTGCTGTGTGAAGAC CTCCCAGGCACGGAAGACTTT	SEQ ID 20 : VFQEAQRQLQKIDKSEGRFHVQN LSQVEQDGRGTGHLRRSSKFCLK EHKALKTLGIIMGTFTLCWLFFIV NIVHVIQDNLIRKEVYILLNWGYV NSGFNPLIYCRSPDFRIAFQELLCL RRSSLKAYGNGYSSNGNTGEQS GYHVEQEKENKLLCEDLPGTEDF VGHQGTVPSDNIDSQGRNCSTND SLL	#	#	#	#
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	GTGGCCCATCAAGGTACTGTG CCTAGCGATAACATTGATTAC AAGGAGGAAATTGTAGTACAAA TGACTCACTGCTATAA					
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hSHP2_FL	SEQ ID 21:	SEQ ID 22:	#	#	#
<p>ATGACATCGCGGAGATGGTTTCACCCAAATA TCACTGGTGTGGAGGCAGAAAACCTACTGTT GACAAGAGGAGTTAATGCCAGTTTTTGGCA AGGCCTAGTAAAGTAACCCCTGGAGACTTCA CACTTTCCGTTAGAAGAAATGGAGCTGTAC CCACATCAAGATTCAGAACACTGGTGATTAC TATGACCTGTATGGAGGGGAGAAATTTGCCA CTTTGGCTGAGTTGGTCCAGTATTACATGGA ACATCAGGGCAATTAAGAGAGAAATGGA GATGCTATTGAGCTTAAATATCCTCTGAAC TG TGCAGATCCTACCTCTGAAAGGTGGTTTCAT GGACATCTCTCTGGGAAAGAGCAGAGAAAT TATTAACTGAAAAGGAAACATGGTAGTTTT CTTGACGAGAGAGCCAGAGCCACCCTGGA GATTTGTCTTCTGTGCGCACTGGTGATG ACAAAGGGAGAGCAATGACGGCAAGTCTA AAGTGACCCATGTTATGATTCGCTGTCAGGA ACTGAAATACGACGTTGGTGAGGAGAACG GTTTGATTCTTTGACAGATCTTGTTGGAACATT ATAAGAAGAAATCCTATGGTGGAACATTGGG TACAGTACTACAACTCAAGCAGCCCTTAAC ACGACTCGTATAAATGCTGCTGAAATAGAAA GCAGAGTTCGAGAACTAAGCAAATTAGCTGA GACCACAGATAAAGTCAACAAGGCTTTTGG GAAGAAATTTGAGACACTACAACAACAGGAGT GCAAACCTCTCTACAGCCGAAAAGAGGGTCA AAGGCAAGAAAACAAAAAATAAGATAT AAAAACATCCTGCCCTTTGATCATACCAAGG TTGTCCTACACGATGGTGATCCCAATGAGCC TGTTTCAGATTACATCAATGCAAAATATCATCA TGCCTGAATTTGAAACCAAGTGCAACAATTC AAAGCCCCAAAAGAGTTACATTGCCACACAA GGCTGCCTGCAAAACACCG</p>	<p>MTSRWFHPNITGVEAENLLTRGVNGS FLARPSKNPGDFTLSVRRNGAVTHIKIQ NTGDYYDLYGGEKFATLAEVLVQYMEH HGQLEKNQDVIELKYPNLCADPTSERW FHGLSGKEAEKLLTEKGKHGSLVRES QSHPGDFVLSVRTGDDKGESNDGKSKV THVMIRCQELKYDVGGGERFDSLTLVE HYKKNPMVETLGTVLQLKQPLNTRINA AEIESRVRELSKLAETTDKVKGWFEEF ETLQQECKLLYSRKEGQRQENKNKNR YKNILPFDHTRVWLHDGDPNEPVS DYNA NIIMPEFETKCNSKPKKSYIATQGCLQN TVNDFWRMVFQENSRVIMTTKEVERG KSKCVKYWPDEYALKEYGVMRVNRVKE SAAHDYTLRELKLSKVGQALLQGNTERT VWQYHFRTPDHPGVSPDPGGVDFLEE VHHQESIMDAGPVVHCSAGIGRTGTF IVIDLIDIREKGVDCDIDVPKTIQMVR SQ RSGMVQTEAQYRFIYMAVQHYIETLQRR IEEEQSKRKGHEYNIKYSLADQTSGD QSPLPCTPTPPCAEMREDSARVYENV GLMQQKKSFR</p>				

Human Melatonin 1a receptor	SEQ ID 27 : GGCATCGCCATCAACCGCTACTGCTACATCT GCCACAGTCTCAAGTACGACAACTGTACAG CAGCAAGAACTCCCTCTGCTACTAG	SEQ ID 28 : GIAINRYCYICHSLKYDKLYSSKNSLCY	#	#	#	#
Human Melatonin 1a receptor	SEQ ID 29 : CTGGTTCTCCAGGTCAGACAGAGGGTGAAA CCTGACCGCAAAACCCAACTGAAACCCACAG GACTTCAGGAATTTGTCACCATGTTTTAG	SEQ ID 30 : LVLQVRQVRVKPDRKPKLKPQDFRNFVT MF	#	#	#	#
Human Melatonin 1a receptor	SEQ ID 31 : TACGGGCTACTGAACCAAAATTTCAGGAAGG AATACAGGAGAATTATAGTCTCGCTCTGTAC AGCCAGGGTGTTCTTTGTGGACAGCTCTAAC GACGTGGCCGATAGGGTTAAATGGAAACCG TCTCCACTGATGACCAACAATAATGTAGTAA GGTGGACTCCGTTTAA	SEQ ID 32 : YGLLNQNFKEYRRIIVSLCTARVFFVDS SNDVADRVKWKPSPLMTNNNNV/KVDSV	#	#	#	#

Human melatonin 1b receptor	SEQ ID 33 : GCCATCGCCATTAAACCGCTACTGCTACATCT GCCACAGCATGGCCTACCAACCGAATCTACC GGCGCTGGCACACCCCTCTGCACTGA	SEQ ID 34 : AIAINRYCYICHSMAYHRIYRRWHTPLH	#	#	#	#
Human melatonin 1b receptor	SEQ ID 35 : CTGGTGCTTCAGGCCCGCAGGAAAGCCCAAG CCAGAGAGCAGGCTGTGCCTGAAGCCCAGC GACTTGGGGAGCTTTCTAACCATGTTTTGA	SEQ ID 36 : LVLQARRKAKPESRLCLKPSDLRSFLTM F	#	#	#	#
Human melatonin 1b receptor	SEQ ID 37 : TATGGGCTCTTGAACCAAACTCCGCAGGG AATACAAGAGGATCCTCTTGGCCCTTTGGAA CCCACGGCACTGCATTCAAGATGCTTCCAAG GGCAGCCACGCGGGGCTGCAGAGCCCC AGCTCCACCCATCATTTGGTGTGCAGCACCAG GCAGATGCTCTCTAG	SEQ ID 38 : YGLLNQNFREYKRILLALWNPRHCIQD ASKGSHAEGLQSPAPPIGVQHQADAL	#	#	#	#

Human OB-receptor short form	SEQ ID 39 : GGAACATTATTAATATACACACCAAGAATGAA AAAGCTATTTGGGAAGATGTTCCGAACCCC AAGAAATTGTTCTGCGCACAAAGGACTTAATT TTCAGAAGAGAGAACGGACATTCTTTGA	SEQ ID 40 : GTLISHQRMKKLFWEDVNPKNCSWA QGLNFQKRTDIL	#	#	#	#
hOB-receptor long form	SEQ ID 41 : GGAACATTATTAATATACACACCAAGAATGAA AAAGCTATTTGGGAAGATGTTCCGAACCCC AAGAAATTGTTCTGCGCACAAAGGACTTAATT TTCAGAAGCCAGAAACGTTTGAGCATCTTTT ATCAAGCATACAGCATCAGTGACATGTGGTC CTCTCTTTTGGAGCCTGAAACAAATTCAGAA GATATCAGTGTTGATACATCATGGAATAA AGATGAGATGATGCCAACAACTGTGGTCTCT CTACTTTCAACACAGATCTTGAAAAGGGTT CTGTTTGATTAGTGACCACTTCAACAGTGT AACTTCTCTGAGGCTGAGGGTACTGAGGTAA CCTATGAGGACGAAAGCCAGAGACAACCCCT TGTTAAATACGCCACGCTGATCTAG	SEQ ID 42 : GTLISHQRMKKLFWEDVNPKNCSWA QGLNFQKPEFHEHLFIKHTASVTCGPLLL EPETISEDISVDTSWKNKDEMMPTTVS LLSTTDLEKGSVCISDQFNVSFSEAEGT EVTYEDESQRQPFVKYATLI	#	#	#	#

hOB-receptor long form	SEQ ID 43 : GCCACGCTGATCAGCAACTCTAAACCAAGTG AAACTGGTGAAGAACAAAGGCTTATAAATAG TTCAGTCACCAAGTGCTTCTCTAGCAAAAATT CTCCATTGAAGGATTCTTCTCTAATAGCTCA TGGGAGATAGAGGCCAGGCATTTTATAT TATCAGATCAGCATCCCAACATAATTTACCCA CACCTCACATTCTCAGAAAGGATTGGATGAAC TTTTGAAATTGGAGGGAATTTCCCTGAAGA AAATAATGATAAAAAGTCTATCTATTATTTAG GGTCACCTCAATCAAAAAGAGAGAGAGTG GTGTGCTTTTGACTGACAAAGTCAAGGGTATC GTGCCCATTCCCAGCCCCCTGTTTATTCACG GACATCAGAGTTCTCCAGGACAGTTGCTCAC ACTTTGTAGAAAATAATCAACTTAGGAAC TCTAGTAAGAAGACTTTTGCATCTTACATGCC TCAATTCCAAAGTTGTTCTACTCAGACTCATA AGATCATGGAAAACAAGATGTGTGACCTAAC TGTGTAA	SEQ ID 44 : ATLISNSKPSSETGEEQGLINSSVTKCFSS KNSPLKDSFSNSSWEIEAQAFFILSDQHP NIISPHLTFSEGLDELLKLEGNFPEENND KKSYYLGVTSIKKRESGVLLTDKSRVSC PPAPCLFTDIRVLQDSCSHFVENNINLG TSSKKTFASYMPPQFQTCSTQTHKIMENK MCDLTV	#	#	#	#
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hOB-receptor long form	SEQ ID 45 : GGAACATTATTAATATACACACCAAGAATGAA AAAGCTATTTGGGAAGATGTTCCGAACCCC AAGAAATGTTCTGGGCACAGGACTTAATT TTCAGAGCCAGAAACGTTTGAGCATCTTTT ATCAAGCATACAGCATCAGTGACATGTGGTC CTCTCTTTGGAGCCTGAACAAATTCAGAA GATATCAGTGTGATACATCATGAAAAATAA AGATGAGATGATGCCAACAACTGTGGTCTCT CTACTTTCAACACAGATCTTGAAAAGGTT CTGTTTGTATTAGTGACCAGTTCAACAGTGT AACTTCTCTGAGGCTGAGGGTACTGAGGTAA CCTATGAGGACGAAAGCCAGAGACAACCCCT TGTTAAATACGCCACGCTGATCAGCAACTCT AAACCAAGTGAAACTGGTGAAGAACAAAGG CTTATAATAGTTCAGTCACCAAGTGCTTCTC TAGCAAAATTCCTCATTGAAGGATCTTTCT CTAATAGCTCATGGGAGATAGAGGCCCAGG CATTTTATATTATCAGATCAGCATCCCAAC ATAATTCACCACACCTCACATTCTCAGAA GATTGGATGAACTTTGAAATTGGAGGAA TTCCCTGAAGAAATAATGATAAAAGTCTA TCTATTATTAGGGTCACCTCAATCAAAAG AGAGAGATGGTGCTTTTGACTGACAAAGT CAAGGGTATCGTGCCCATTCAGAGCCCT GTTTATCAGGACATCAGAGTCTCCAGGA CAGTTGCTCACACTTTGAGAAAATAATATCA ACTTAGGAACCTTAGTAAGAAGACTTTTGCA TCTTACATGCCTCAATCCAACTTGTTCTAC TCAGACTCATAAGATCATGGAACAAAGATG TGTGACCTAACTGTGTA	SEQ ID 46 : GTLISHQRMKKLFWEDVPNPKNCWSA QGLNFQKPETFEHLFIKHTASVTCGPLLL EPETISEDISVDTSWKNKDEMMPTTVS LLSTTDLEKGSVCISDQFNSVNFSEAGT EVTYEDESQRQPFVKYATLISNSKPSET GEEQGLINSSVTKCFSSKN SPLKDSFSN SSWEIEAQAFFILSDQHPNISPHTFSEG LDELLKLEGNFPEENNDDKSIYYLGVTSI KKRESGVLLTDKSRVSCFPFAPCLFTDIR VLQDSCSHFVENNINLGTSSKKTFSYM PQFQTCSTQTHKIMENKMCDLTV	#	#	#	#
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Human ADBR kinase 1	SEQ ID 47 :	SEQ ID 48 :	#	#	#
ATGGCGGACCTGGAGGCGGTGCTGGCCGA CGTGAGCTACCTGATGGCCATGGAGAAGAG CAAGCCACGCGCGCGCGCGCGCCAGCA AGAAGATACTGCTGCCCGAGCCAGCATCC GCAGTGTATGCAGAAGTACCTGGAGGACC GGGCGAGGTGACCTTTGAGAAGATCTTTTC CCAGAAGCTGGGTACCTGCTCTCCGAGA CTTCTGCTGAACCACTGGAGGAGGCCAG GCCCTGGTGGAA TCTATGAGGAGATCAAG AAGTACGAGAAGCTGGAGCGGAGGAGGAG CGTGTGCGCGCAGCGCGGAGATCTTCGAC TCATACATCATGAAGGAGCTGCTGGCCTGCT CGCATCCCTTCTCGAAGAGTGCCACTGAGCA TGTCGAAGGCCACCTGGGAAGAAGCAGGT GCCTCCGGATCTCTCCAGCCATACATCGAA GAGATTTGTCAAAACCTCCGAGGGACGTGT TCCAGAAATTCATTGAGCGGATAAGTTAC ACGGTTTGGCCAGTGGAAGAAATGTGGAGCTC AACATCCACCTGACCATGAATGACTTCAGCG TGCATCGCATCATTTGGCGCGGGGCTTTG GCGAGGTCTATGGTGCCGGAAGGCTGACA CAGGCAAGATGTACGCCATGAAGTGCCTGG ACAAAAAGCGCATCAAGATGAAGCAGGGGG AGACCTGGCCCTGAACGAGCGCATCATGC TCTCGCTCGTCAGCACTGGGACTGCCCAT CATTGTCTGCATGTACACGCGTTCACACG CCAGACAAGCTCAGCTTCACTCGACCTCA TGAACGGTGGGACCTGCACTACCACTCT CCCAGCAGGGGTCTTCTCAGAGGCTGACA TGCGCTTCTATGGCGCGGAGATCATCCTGG GCCTGGAGCACATGCACAACCGCTTCGTGG TCTACCGGACCTGAAGCCAGCCCAACATCCT TCTGGACGAGCATGGCCACGTCGGGATCTC GGACCTGGGCC	MADLEAVLADVSYLMAMEKSKATPAARA SKILLPEPSIRSVMQYLEDRGEVTFEKI FSQKLGYLFRDFCLNHLEEARPLVEFYE EIKYKLETEEEERVARSEIFDSYIMKEL LACSHPFKSATEHVQHLGKKQVPPDL FQPYEEICQNLRGDVQKFIESDKFTRF CQWKNVELNIHLTMNDFS VHRIGRGF GEVYGRKADTGKMYAMKCLDKRIKM KQGETLALNERIMLSLVSTGDCPFIVCMS YAFHTPDKLSFILDLMNGGDLHYHLSQH GVFSEADMRFYAAEILGLEHMHNRFW YRDLK PANILLDEHGHVRISDLGLACDFS KKKPHASVGTGMYAPEVLQKGVAYDS SADWFSLGCMLFKLLRGHSPFRQHKTK DKHEIDRM TLTMAVELPDSFSPELRSLE GLLRQDVNRRLGCLGRGAQEVKESPF RSLDWQMVFLOKYPPLIPPRGEVNAAD AFDIGSFDEEDTKGKLLDSQDEL YRNFP LTISERWQVEAETVFD TNAETDRLEAR KKAKNKQLGHEEDYALGKDCIMHGYS KMGNPFLTQWQRRYFYLPNRLWRGE GEAPQSLTMEEQSVETQIKERKCLLL KIRGGKQFILQCDSDEL VQWKELRDA YREAQQLVQRVPMKNKPRSPVELSK VPLVQRGSANGL				

Rat ADBR kinase 2	SEQ ID 49 : ATGGCGGACCTGGAGGCGCGTGTGGCCGAT GTCAGTTACCTGATGGCCATGGAGAAGAGC AAGGCGACCCCGCGCGCCGCGCCAGCAA GAGGATCGTCTCGCGGAGCCAGTATCCG GAGTGTGATGCAGAAGTACCTTGCAGAGAG AAATGAAATAACCTTTGACAAGATTTTCAATC AGAAAATTGGTTCTTGCTATTTAAAGATTTT TGTTTGAATGAAATTAATGAAGCTGTACCTCA GGTGAAGTTTATGAAGAGATAAAGGAATAT GAAAACCTTGATAATGAGGAAGACCGCTTT GCAGAAGTCGACAAATTTATGATGCCTACAT CATGAAGGAACCTTCTTCTGTTCCATCCTT TCTCAAAGCAAGCTGTAGAACACGTACAAG TCATTTATCCAAGAAACAAGTGACATCAACTC TTTTTCAGCCATACATAGAAGAAATTTGTAA AGCCTTCGAGGTGACATTTTCAAAAATTTAT GGAAAGTGACAAGTTCAGTAGATTTGTGAG TGGAAAACGTTGAATTAATATCCATTTGAG CATGAATGAGTTCAGTGTGATAGGATATT GGACGAGGAGGATTCGGGGAAGTTTATGGT TGCAGGAAAGCAGACACTGGAAAAATGTATG CAATGAAATGCTTAGATAAGAAAGAGGATCAA AATGAAACAAGGAGAAACATTAGCCTTAAAT GAAAGAATCATGTTGTCTCTTGTGACACAG GAGACTGTCCTTTTCATTGTATGTATGACCTAT GCCTTCCATACCCAGATAAACTCTGCTTCA TCCTGGATCTGATGAACGGGGCGGATTTC ACTACCACCTTTCAACACACGGTGTGTTCTC TGAGAAGGAGATGCGGTTTTTATGCCACTGAA ATCATTCTGGGTCTGGAACACATGCACAATC GGTTTGTGCTACAGAGATTTGAAGCCAGC AAATATTCTCTTGGATGAACATGGACACGCA AGAATATCAGATCTTGGTC	SEQ ID 50 : MADLEAVLADVSYLMAMEKSKATPAARA SKRIVLPEPSIRSMQKYLAEARNEITFDKI FNQIGIFLLFKDFCLNEINEAVPQVKFYE EIKEYEKLDNEEDRLCRSRQIYDAYIMKE LLSCSHPFKQAVEHVQSHLSKKQVTST LFQPYEEICESLRGDIQKFMESEDKFTR FCQWKNVELNIHLMNEFSVHRIIGRGG FGEVYGCRAKADTGKMYAMKCLDKKRIK MKQGETLALNERIMLSLVSTGDCPFIVC MTYAFHTPDKLCFILDLMNGGDLHYHLS QHGVFSEKEMRFYATEILGLEHMHNR VYVRDLKPANILLDEGHARISDLGLACD FSKKKPHASVGTGMYMAPEVLQKGTAY DSSADWFSLGCMLFKLLRGHSPFRQHK TKDKHEIDRMTLTVNVELPDTFSPELKS LEGLQRDVSKRLGCHGGSGQEVKEHS FFKGVDWQHVYLQKYPPLIPPRGEVNA ADAFDIGSFEEDTKGIKLLDCDQELYKN FPLVISERWQQEVETVYEAVNADTDKIE ARKRAKNQQLGHEEDYALGKDCIMHGY MLKLGNPFLTQWQRRYFYLFPNRLEWR GEGESRQNLMTMEQILSVEETQIKDKKCI LFRIKGGKQFVLQCESDPEFVQWKKEIN ETFKEAQRLLRRAPKFLNKPRSGTVELP KPSLCHRNSSGL	#	#	#	#
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Rat beta Arrestin 1	<p>SEQ ID 51 :</p> <p>ATGGCGACAAAGGGACGCGGGTGTCAAG AAGCGAGCCCCAATGGAAGCTCACCGTC TATCTGGGAAAGCGGACCTTTGTGGACCACA TCGACCTCGTGGAGCCCGTGGATGGAGTGG TTCTTGTGATCCGAGTATCTCAAGGAGAG GAGAGTCTATGTACGCTGACCTGCGCCTTC CGCTACGGCCGGAGGACCTGGATGTCCTG GGCCTGACCTTCGCAAGGACCTGTTGTGG CCAACGTGCAGTCTTCCCGCGGCCCTG AGGACAAGAAGCCCTGACGCGGCTGCAGG AGCGCTCATCAAGAAGCTGGCGGAGCATG CCTACCCCTTTCACCTTTGAGATCCCTCCGAA CCTCCCATGCTCTGTGACTTGCAGCCGGGA CCTGAAGATACAGGGAAGCCTGCGGTGTG GACTACGAAGTGAAGCCTTCTGTGCGGAG AACCTGGAGGAGAAGATCCACAAGCGGAAT TCTGTGCGCCTGGTCTATCCGGAAGTTTCAGT ATGCCCCAGAGAGGCTGCCCCCAGCCGA CGCCGAGACCAAGGAGCTTCTCATGT CAGACAAGCCCTTGCATCTGGAGCCTCCC TGGACAAGGAGATCTACTACCACGGAGAAC CCATCAGTGTCAACGTCCATGTCAACCAAA CACCAACAAGACGGTGAAGAAGATCAAGATC TCGGTGGCCAGTATGCAGACATCTGTCTGT TCAACACAGCCCGAGTACAAGTCCCTGTGG CCATGGAAGAGGCTGATGACACAGTGGCAC CCAGCTCTACGTTCTGCAAGGTCTACACGCT GACCCCTTCTGGCCAACAATCGAGAGAA GCGGGGCTCGCCCTGGACGGGAAGCTCAA ACACGAGGACACGAACCTGGCCTCCAGCAC CCTGTTGAGGGAAGGAGCCAAACCGGGAGAT CCTGGGCATCATGTTTCTCTACAAAGTGAAA GTGAAGCTGGTGGTGTCTCGTGGCGGCGCTG TTGGGAGATC</p>	<p>SEQ ID 52 :</p> <p>MGDKGTRVFKKASPNGLTVYLGKRD VDHIDLVEPVDGWLVDP EYLKERRVYV TLTCAFYRGREDLDVLGLTRKDLFVAN VQSFPPAPEDKKPLTRLQERLIKLG YPTFEIPPNLPCSVTLQPGEDTGKAC GVDYEVKAFCAENLEEKHKRNSVRLVIR KVQYAPERPGPQPTAETTRQFLMSDKPL HLEASLDKEIYYHGEPISVNVHVTNNTNK TVKKISVRQYADICLFNTAQYKCPVAM EEADDTVAPSSTFCKVYTLTPFLANNRE KRGALDGLKHEDTNLASSTLLREGAN REILGIIVSYKVVKLVVSRGGLGDLASS DVAVELPFTLMHPKPEEPHREVPEHE TPVDTNLIELTDNDDIVFEDFARQLKG MKDDKEEEDGTGSPRTRELRSMSLL VVLLWNYWKVRNCQVLLSYPKRNKLN</p>	#	#	#	#
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Rat beta aArestin2	SEQ ID 53 : ATGGTGAAAAACCCGGGACCAAGGCTCTC AAGAAGTCGAGCCCTAACTGCAAGCTCACC GTGTACTTGGCAAGCGTGACTTTGTGGATC ACTTGGACAAAGTGATCCTGTCGATGGTGT GGTCTTGTGGATCCTGACTACTTGAAGGAC CGGAAAGTGTGTGACCCCTACCTGTGCCT TCCGCTATGGCGGAGAACCTGGATGTACT GGCCCTGTCTTCCGCAAGATCTGTTTCATC GCCACCTACGAGCCTTCCCCCATGCC AACCCACCTCGGCCCCACCCGCTACAG GACCGACTGCTGAAGAAGTTGGGCCAGCAT GCCACCCCTTTTTTTCACAATACCCGAGA ATTGCTTGTCTCCGTCACACTGCAGCCAGG ACCGGAGGACACAGGGAAGGCTGTGGAGT AGACTTTGAGATTCGAGCCTTCTGTGCCAA TCTATAGAAAAAAGCCACAAAAGGAAC CCGTGCGGCTTATCATCAGAAAGGTACAGTT TGCTCTGAGACACCCGCCCCCAGCCATC AGCTGAAACCCACACGCCACTTCCTCATGTCT GACCGAGGTCCCTGCACCTAGAGGCTTCC CTGGACAAAGAGCTGTACTACCATGGGAA CCCCTCAATGTCAAGTCCAGGTACCAACA ATTCTGCCAAGACCGTCAAGAAGATCAGAGT GTCTGTGAGACAGTATGCCGACATTTGCCCTC TTCAGCACCGCGCAGTACAAGTGTCTGTG GCTCAGCTTGAACAAGATGACCAAGGTGTCTC CCAGTCCACATTCTGCAAGGTGTACACCAT AACCCCGCTGCTCAGTGACAAACCGAGAGAA GCGTGGCTTGGCTTGTGAGGCAACTCAA GCACGAAGACACCAACCTGGCTTCCAGCAC CATTGTGAAGGAGGAGGCCAACAAAGGAGT GCTGGGAATCCTAGTATCCTACAGGCTCAAG GTGAAGCTGGTGTCTCGAGGCGGGGAT GTCT	SEQ ID 54 : MGEKPGTRVFKKSSPNCKLTVYLGRDF VDHLDKVDVPDGVWLVDPDYLKDRKVFEV TLTCAFRYGREDLDVLGLSFRKDLFIATY QAFPPMPNPPPTRLQDRLLKLLGQH AHPFFFTIPQNLPCSVTLQPGPEDITGKA CGVDFEIRAFCAKSEIEKSHKRNSVRLIIR KVQFAPETPGPQPSAETTRHFLMSDRR SLHLEASLDKELYHGEPLNVNVHVTNN SAKTVKKIRSVRQYADICLFTAQYKCP VAQLEQDDQVSPSSTFCKVYTIITLLSDN REKRLALDGGQLKHEDTNLASSTVKEG ANKEVLGILVSYRVKVKLVVSRGGDVSV ELPFVLMHPKPHDHITLPRPQSAPREIDI PVDNLIEFDNTYATDDDIVFEDFARLRL KGMKDDDCDDQFC	#	#	#	#
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human STAT3 : Transcription factor; Phosphorylation by JAK-type kinases leads to dimerisation and translocation to the nucleus to transactivate target gene expression	SEQ ID 55 : ATGCCCAATGGAATCAGCTACAGCAGCTTG ACACAGGTACCTGGAGCAGCTCCATCAGC TCTACAGTGACAGCTTCCCAATGAGCTGCG GCAGTTCTGCCCCCTTGATTGAGAGTCAA GATTGGCATAATGCGGCCAGCAAGAATCA CATGCCACTTTGGTGTTCATAATCTCCTGG GAGAGATTGACCAGCAGTATAGCGCTTCCT GCAAGAGTCGAATGTTCTCTATCAGCACAAT CTACGAAGAATCAAGCAGTTTCTTCAAGACA GGTATCTGGAGAAGCCAATGGAGATTGCC GGATTGTGCCCGGTGCTGTGGGAAGAAT CACGCTTCTACAGACTGCAGCCACTGCGG CCCAGCAAGGGGCCAGGCCAACCCACCCCA CAGCAGCCGTGGTGACGGAGAAAGCAGACA TGCTGGAGCAGCACCTTCAGGATGTCCGGA AGAGAGTGCAGGATCTAGAACAGAAAATGAA AGTGGTAGAGAACTCCAGGATGACTTTGAT TTCAACTATAAACCTCAAGAGTCAAGGAG ACATGCAAGATCTGAATGGAAACAACCACTC AGTGACCGAGCAGAAAGATGCAGCAGCTGGA ACAGATGCTCACTGCGCTGGACCAGATGCG GAGAAGCATCGTGAGTGAGCTGCGCGGGCT TTTGTCAGCGATGGAGTACGTGCAGAAAAC CTACGGACGAGGAGCTGGCTGACTGGAAG AGCGGCAACAGATTGCTGCTGATTGGAGGC CCGCCAACATCTGCCCTAGATCGGCTAGAAA ACTGATAACGTCAATAGCAGAACTCAACT TCAGACCCGTCACAAATTAAGAACTGGAG GAGTTGCAGCAAAAAGTTTCTACAAAGGGG ACCCATTGTACAGCACCGGCCGATGCTGG AGGAGAGAAATCGTGAGCTGTTTAGAACTT AATGAAAAGTGCCCTTTGTGGTGAGCGGCA GCCCTGCATGCCCCATGCATCCTGACCCGGC CCTCG	SEQ ID 56 : MAQWNQLQLDTRYLEQLHQLYSDSFP MELRQLAPWIESQDWAYAAKESHATL VFHLLGEIDQQYSRFLQESNVLYQHNL RRIKFLQSRYLEKPMELRIVARCLWEE SRLQTAATAAQGGQANHPATAAWE KQMLEQLHLDVRKRVQDLEQKMKVVE NLQDDFFNYKTLKSQDMQDLNGNNQ SVTRQKMQLEQMLTALDQMRRSIVSEL AGLLSAMEYVQKTLTDEELADWKRRQOI ACIGPPNICLDRLNENWITSLAESQLQTR QKIKLEELQQKVSYGDPVQHRPML ERIVELFRNLMSAFWERQPCMPMHPD RPLVIKTGVQFTTKVRLLVKFPENYQLKI KVCIDKSDGVAALRGSRKFNILGTNTKV MNMEESNNGSLSAEFKHLTLREQRCGN GGRANCDASLIVTEELHLITFETEVYHQG LKIDLETHSLPWVWISNICMPNAWASIL WYNMLTNPNKNVFFTKPIGTWDQVA EVLSWQFSSTTKRGLSIEQLTTLAEKLLG PGVNYSGCQITWAKFCKENMAGKGFSE WWWLDNIIDLVKYILALWNEGIMGFISK ERERAILSTKPPGTFLRFSESSKEGGVT FTWVEKDISGKTQIQSVEPYTKQQLNNM SFAKIMGYKIMDATNILVSPLVLYPDIPK EEAFGKYCRPESQEHPEADPGSAAPYL KTKFCVTPTTCSNTIDLPMSPRTLDSLM QFGNNGEGAEPGAGGQFESLTFDMELT SECATSPM	#	#	#	#
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mCIS	SEQ ID 57 :	SEQ ID 58 :	#	#	#	#
	<p>ATGGTCTCTGCGTACAGGATCTTGTCCTT TGCTGGCTGTGGAGCAAAATTGGCGGCGGC CTCTGTGGCCAGTCCCTGGAGCTGCCCG GGCAGCCATGCAGCCCTACCCACTGGGG CATCCAGAGGAAGTGACAGAGGAGACCC CTGTCCAGGCAGAGAAATGAACCGAAGGTGC TAGACCTGAGGGGATCTGCTGTGCATAG CCAAGACGTTCTCCTACCTTCGGGAATCTGG GTGGTACTGGGTTCTATTACAGCCAGCGA GGCCCGGCAGCACCTACAGAAGATGCCGGA GGGTACATTCTAGTTCGAGACAGCACCCAC CCCAGCTACCTGTTACACACTGTCAAGTCAAAA CCACCGGTGGCCCAACCAACGTGCGGATCG AGTACGCCGATTCTAGCTTCGGGTGGACTC TAAGTCTGTGCAAGACCTCGAATCCTGGCC TTCCCAGATGTGGTCAGCCTTGTCAGCACT ATGTGGCTCCTGTGCAGCTGACACCCGGA GCGACAGCCCGATCCTGCTCCCAACCCAG CCCTGCCTATGCTAAGCAAGATGCACCTAG TGACTCGGTGCTGCCTATCCCGTGGCTACT GCAGTGCACTGAAACTGGTGACGCCCTTT GTGCGCAGGAGAGTGCCCGCAGCTTACAA CATCTGTGCGGCTAGTCATCAACCGTCTGG TGGCCGACGTGGACTGCTTACCCCTGCCCC GGCGTATGGCCGACTACCTCCGACAGTACC CCTTCCAACCTCTGA</p>	<p>MVLCVQGSCLLAVEQIGRRPLWAQSLE LPGPAMQPLPTGAFPEEVTEETPVQAEN EPKVLDPEDLLCIAKTFSYLRESGWYW GSITASEARQHLQKMEGTFLVRDSTHP SYLFTLSVKTTTRGPTNVRIEYADSSFRLD SNCLSRPRILAFPDVWSLVQHYVASCAA DTRSDSPDPAPTALPMSKQDAPSDSVL PIPVATAVHLKLVQPFVRRSSARSLOHLC RLVINRLVADVDCCLPRRMADYLRQYP FQL</p>				

mSOCS1	<p>SEQ ID 59 :</p> <p>ATGGTAGCACGCAACACGAGGTGGCAGCCGAC AATGCGATCTCCCGGAGCAGAGCCCCGA CGGCGTGACAGCCCTCCTCGCTCGTCT TCGTCTCGCCAGCGGCCCCGTCGTCCTC CGGCCCTGCCGGGCTCCAGCCCCAGC CCCTGGCGACACTCACTCCGCACCTTCCG CTCCCACTCCGATTACCGGCGCATCACGCG GACCAGCGCGCTCCTGGACGCTGCGGCTT CTATTGGGACCCCTGAGCGTGCACGGGGC GCACGAGCGGCTGCGTGCCGAGCCCGTGG GCACCTTCTTGGTGCGCGACAGTCGCCAAC GGAAGTCTTCTTCGCGCTCAGCGTGAAGAT GGCTCGGGCCCCACGAGCATCCGCGTGCA CTCCAGGCGGCGCTTCCACTTGGACGG CAGCCGCGAGACCTTCGACTGCCTTTTCA GCTGCTGGAGCACTACGTGGCGGCGCGCG GCCGATGTTGGGGCCCCGCTGCGCCAG CGCCGCGTGCGGCGCTGCAGGAGCTGTGT CGCCAGCGCATCGTGCGGCGCGTGGGTG CGAGAACCTGGCGCGCATCCCTCTTAACCC GGTACTCCGTGACTACCTGAGTTCTTCCCC TTCAGATCTGA</p>	<p>SEQ ID 60 :</p> <p>MVARNQVAADNAISPAAEPRRRSEPSSS SSSSSPAAPVRPRPCPAVPAPAGDTHF RTFRSHDYRRITRTSALLDACGFYWG LSVHGAHERLRAEPVGTFLVRDSRQRN CFFALSVKMASGPTSIRVHFQAGRFHLD GSRETFDCLFELLEHYVAAPRMLGAPL RQRRVRPLQELCRQRIAAVAVGRENLARI PLNPVLRDYLSSFPEQI</p>	#	#	#	#
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Table 2: bait-prey interactions

1-Bait Protein Name	2-Amino acid sequence	3-Prey Gene Name	
Human Skp1	SEQ ID 2	gb AB007952 AB007952 Homo sapiens mRNA for KIAA0483 protein, partial cds.	
Human Skp1	SEQ ID 2	gb AB019987 AB019987 Homo sapiens mRNA for chromosome-associated polypeptide-C, complete cds.	
Human Skp1	SEQ ID 2	gb AB033279 AB033279 Homo sapiens BTRCP2 mRNA for F-box and WD-repeats protein beta-TRCP2 isoform A, complete cds.	
Human Skp1	SEQ ID 2	gb AF011751 AF011751 Hepatitis C virus strain H77 pCV-H77C polyprotein gene, complete cds.	
Human Skp1	SEQ ID 2	gb AF054284 AF054284 Homo sapiens spliceosomal protein SAP 155 mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AF129534 AF129534 Homo sapiens chromosome 5 F-box protein Fbx4 (FBX4) mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AF142481 AF142481 Homo sapiens F-box protein FLR1 (FLR1) mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AF157323 AF157323 Homo sapiens p45SKP2-like protein mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AF174599 AF174599 Homo sapiens F-box protein Fbx11 (FBX11) mRNA, partial cds.	
Human Skp1	SEQ ID 2	gb AF176698 AF176698 Homo sapiens F-box protein FBW2 mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AF179221 AF179221 Homo sapiens F-box protein Liliina (LILINA) mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AF184275 AF184275 Mus musculus F-box protein FBX18 mRNA, partial cds.	
Human Skp1	SEQ ID 2	gb AF199356 AF199356 Homo sapiens F-box protein FBL6 (FBL6) mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AF233225 AF233225 Homo sapiens F-box protein FBX (FBX) mRNA, complete cds.	
Human Skp1	SEQ ID 2	gb AK001933 AK001933 Homo sapiens cDNA FLJ11071 fis, clone PLACE1004937, moderately similar to SEL-10 PROTEIN.	
Human Skp1	SEQ ID 2	gb AL034374 HS483K16 Human DNA sequence from clone RP3-483K16 on chromosome 6p12.1-21.1.	
Human Skp1	SEQ ID 2	gb AL109627 HSJ733M16 Human DNA sequence from clone RP4-733M16 on chromosome 1p36.11-36.23, complete sequence.	
Human Skp1	SEQ ID 2	gb D25542 HUMGCP372 Human mRNA for golgi antigen gcp372, complete cds.	
Human Skp1	SEQ ID 2	gb D29954 HUMORFA06 Human mRNA for KIAA0056 gene, partial cds.	
Human Skp1	SEQ ID 2	gb G23579 G23579 human STS WI-15073, sequence tagged site.	
Human Skp1	SEQ ID 2	gb M67463 HPCCGAA Hepatitis C virus, complete genome.	
Human Skp1	SEQ ID 2	gb X17025 HSC54 Human homolog of yeast IPP isomerase.	
Human Splicing Factor 1	SEQ ID 4	gb AB002533 AB002533 Homo sapiens mRNA for Qip1, complete cds.	
Human Splicing Factor 1	SEQ ID 4	gb AB007890 AB007890 Homo sapiens mRNA for KIAA0430 protein, partial cds.	
Human Splicing Factor 1	SEQ ID 4	gb AB011134 AB011134 Homo sapiens mRNA for KIAA0562 protein, complete cds.	

Human Splicing Factor 1	SEQ ID 4	gb AB012190 AB012190 Homo sapiens mRNA for Nedd8-activating enzyme hUba3, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AB020714 AB020714 Homo sapiens mRNA for KIAA0907 protein, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AB032254 AB032254 Homo sapiens BAZ2A mRNA for bromodomain adjacent to zinc finger domain 2A, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AB037781 AB037781 Homo sapiens mRNA for KIAA1360 protein, partial cds.
Human Splicing Factor 1	SEQ ID 4	gb AB037839 AB037839 Homo sapiens mRNA for KIAA1418 protein, partial cds.
Human Splicing Factor 1	SEQ ID 4	gb AC007688 AC007688 Homo sapiens 12p12-27.2-31.7 BAC RPC111-392P7 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
Human Splicing Factor 1	SEQ ID 4	gb AF005361 HUMIMPA6 Homo sapiens importin alpha 6 mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF022770 AF022770 Mus musculus peripheral benzodiazepine receptor associated protein (Pap7) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF029308 HTCRBCHR9 Homo sapiens chromosome 9 duplication of the T cell receptor beta locus and trypsinogen gene families.
Human Splicing Factor 1	SEQ ID 4	gb AF034756 AF034756 Homo sapiens importin-alpha homolog (SRP1gamma) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF038564 AF038564 Homo sapiens atrophin-1 interacting protein 4 (AIP4) mRNA, partial cds.
Human Splicing Factor 1	SEQ ID 4	gb AF046024 AF046024 Homo sapiens UBA3 (UBA3) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF049523 AF049523 Homo sapiens huntingtin-interacting protein HYP4/FBP11 (HYPA) mRNA, partial cds.
Human Splicing Factor 1	SEQ ID 4	gb AF057569 AF057569 Homo sapiens upstream regulatory element binding protein 1 (UREB1) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF060543 AF060543 Homo sapiens importin alpha 7 subunit mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF065485 AF065485 Homo sapiens sorting nexin 4 mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF083243 HSPC025 Homo sapiens HSPC025 mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF111162 AF111162 Homo sapiens guanine nucleotide exchange factor mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF113615 AF113615 Homo sapiens FH1/FH2 domain-containing protein FHOS (FHOS) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF191298 AF191298 Homo sapiens vacuolar sorting protein 35 (VPS35) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb AF200348 AF200348 Homo sapiens melanoma-associated antigen MG50 mRNA, partial cds.

Human Splicing Factor 1	SEQ ID 4	gb AF205588 AF205588 Homo sapiens ZNF01 and HUMORFKG1B genes, partial sequence, complete sequence.
Human Splicing Factor 1	SEQ ID 4	gb AJ010069 HSA010069 Homo sapiens mRNA for HMGBCG protein.
Human Splicing Factor 1	SEQ ID 4	gb AJ242910 HSA242910 Homo sapiens mRNA for N-Acetylglucosamine kinase.
Human Splicing Factor 1	SEQ ID 4	gb AL121973 HSA01012 Human DNA sequence from clone RP3-401O12 on chromosome 6p11.2-21.1 Contains STSs, complete sequence.
Human Splicing Factor 1	SEQ ID 4	gb AL353771 AL353771 Human DNA sequence from clone RP4-677H15 on chromosome 1p31.3-32.3, complete sequence.
Human Splicing Factor 1	SEQ ID 4	gb B48558 B48558 RPCI11-2115.TV RPCI-11 Homo sapiens genomic clone RPCI-11-2115, DNA sequence.
Human Splicing Factor 1	SEQ ID 4	gb D21260 HUMORFEA Human mRNA for KIAA0034 gene, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb E13124 E13124 Human gene for Rho protein-dependent protein kinase, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb L13210 HUMMAC2A Human Mac-2 binding protein mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb L25616 HUMCG1X Homo sapiens kinectin mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb L33075 HUMIQGA Homo sapiens ras GTPase-activating-like protein (IQGAP1) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb M14752 HUMABLA Human c-abl gene, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb M24487 HUMPYHBSB Human prolyl 4-hydroxylase alpha subunit mRNA, complete cds, clone PA-15.
Human Splicing Factor 1	SEQ ID 4	gb M27024 HUMHSP89KD Homo sapiens heat shock protein (HSP89-alpha) gene, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb M35296 HUMARGCAA Human tyrosine kinase arg gene mRNA.
Human Splicing Factor 1	SEQ ID 4	gb M67463 HPCCGAA Hepatitis C virus, complete genome.
Human Splicing Factor 1	SEQ ID 4	gb M75883 HUMSCP2A Human sterol carrier protein X/sterol carrier protein 2 mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb M95178 HUMACTN1A Human non-muscle alpha-actinin mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb S75295 S75295 nucleoprotein interactor 1-SRP1 homolog [human, cervical carcinoma HeLa cells, mRNA, 2940 nt].
Human Splicing Factor 1	SEQ ID 4	gb S78653 S78653 mrg=mas-related [human, Genomic, 2416 nt].
Human Splicing Factor 1	SEQ ID 4	gb U05040 HSU05040 Human FUSE binding protein mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb U07563 HSABLGR3 Human proto-oncogene tyrosine-protein kinase (ABL) gene, exon 1a and exons 2-10, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb U12596 HSU12596 Human tumor necrosis factor type 1 receptor associated protein (TRAP2) mRNA, complete cds.

Human Splicing Factor 1	SEQ ID 4	gb U28386 HSU28386 Human nuclear localization sequence receptor hSRP1alpha mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb U33760 HSU33760 Human cyclin A/CDK2-associated p19 (Skp1) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb U70372 RNU70372 Rattus norvegicus PAM COOH-terminal interactor protein 2 mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb U80213 HSU80213 Human protein arginine N-methyltransferase 2 (PRMT2) mRNA, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb U92704 MMU92704 Mus musculus Olf-1/EBF-like-2(OS) transcription factor (O/E-2(OS)) mRNA, alternative splice variant, complete cds.
Human Splicing Factor 1	SEQ ID 4	gb U96113 HSU96113 Homo sapiens Nedd-4-like ubiquitin-protein ligase WWP1 mRNA, partial cds.
Human Splicing Factor 1	SEQ ID 4	gb X57527 HSCOL8A1 Human COL8A1 mRNA for alpha 1(VIII) collagen.
Human Splicing Factor 1	SEQ ID 4	gb X64044 HSU2AF H.sapiens mmRNA for large subunit of splicing factor U2AF.
Human Splicing Factor 1	SEQ ID 4	gb Y08991 HSP150 H.sapiens mRNA for adaptor protein p150.
mouse p53	SEQ ID 6	gb AB011148 AB011148 Homo sapiens mRNA for KIAA0576 protein, partial cds.
mouse p53	SEQ ID 6	gb AB014530 AB014530 Homo sapiens mRNA for KIAA0630 protein, partial cds.
mouse p53	SEQ ID 6	gb AB021868 AB021868 Homo sapiens PIAS3 mRNA for protein inhibitor of activated STAT3, complete cds.
mouse p53	SEQ ID 6	gb AB035898 AB035898 Homo sapiens hklp2 mRNA for kinesin-like protein 2, complete cds.
mouse p53	SEQ ID 6	gb AF004849 AF004849 Homo sapiens PKY protein kinase mRNA, complete cds.
mouse p53	SEQ ID 6	gb AF060181 AF060181 Homo sapiens zinc finger protein (ZNF198) mRNA, complete cds.
mouse p53	SEQ ID 6	gb AF072825 AF072825 Homo sapiens Raf responsive zinc finger protein (RREB1) mRNA, partial cds.
mouse p53	SEQ ID 6	gb AF077954 AF077954 Homo sapiens protein inhibitor of activated STAT protein PIASx-beta mRNA, complete cds.
mouse p53	SEQ ID 6	gb AF164678 AF164678 Homo sapiens FLASH homolog RIP25 (RIP25) mRNA, complete cds.
mouse p53	SEQ ID 6	gb AF167160 AF167160 Homo sapiens protein inhibitor of activated STAT-1 (PIAS1) mRNA, complete cds.
mouse p53	SEQ ID 6	gb AQ112228 AQ112228 CIT-HSP-2371L12.TR CIT-HSP Homo sapiens genomic clone 2371L12, DNA sequence.
mouse p53	SEQ ID 6	gb AQ487168 AQ487168 RPCI-11-265J2.TV RPCI-11 Homo sapiens genomic clone RPCI-11-265J2, DNA sequence.
mouse p53	SEQ ID 6	gb J03040 HUMSPARC Human SPARC/osteonectin mRNA, complete cds.

mouse p53	SEQ ID 6	gb IM14694 HUMTP53A Human p53 cellular tumor antigen mRNA, complete cds.
mouse p53	SEQ ID 6	gb IM60119 HUMEP2AA Homo sapiens HIV-EP2/Schnurri-2 gene, complete cds.
mouse p53	SEQ ID 6	gb U13843 XXU13843 pBPV cloning vector, complete sequence.
mouse p53	SEQ ID 6	gb U16799 HSU16799 Human Na,K-ATPase beta-1 subunit mRNA, complete cds.
mouse p53	SEQ ID 6	gb U66867 HSU66867 Human ubiquitin conjugating enzyme 9 (hUBC9) mRNA, complete cds.
mouse p53	SEQ ID 6	gb U94788 HSU94788 Human p53 (TP53) gene, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AB007930 AB007930 Homo sapiens mRNA for KIAA0461 peroteine, partial cds.
Human beta-TrCP1	SEQ ID 8	gb AB011089 AB011089 Homo sapiens mRNA for KIAA0517 protein, partial cds.
Human beta-TrCP1	SEQ ID 8	gb AB011148 AB011148 Homo sapiens mRNA for KIAA0576 protein, partial cds.
Human beta-TrCP1	SEQ ID 8	gb AB019002 AB019002 Homo sapiens MRP5 mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AB021868 AB021868 Homo sapiens PIAS3 mRNA for protein inhibitor of activated STAT3, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AB029343 AB029343 Homo sapiens HCR (a-helix coiled-coil rod homologue) gene, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AB037825 AB037825 Homo sapiens mRNA for KIAA1404 protein, partial cds.
Human beta-TrCP1	SEQ ID 8	gb AC005789 AC005789 Homo sapiens chromosome 19, cosmid F5960, complete sequence.
Human beta-TrCP1	SEQ ID 8	gb AF003924 AF003924 Homo sapiens zinc finger protein ANC_2H01 mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF010315 AF010315 Homo sapiens Pig11 (PIG11) mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF060181 AF060181 Homo sapiens zinc finger protein (ZNF198) mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF061836 AF061836 Homo sapiens putative tumor suppressor protein (RDA32) mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF062536 AF062536 Homo sapiens cullin 1 mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF064087 AF064087 Homo sapiens cullin 3 mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF077954 AF077954 Homo sapiens protein inhibitor of activated STAT protein PIASx-beta mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF084940 AF084940 Homo sapiens beta-arrestin 1B mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF116343 AF116343 Homo sapiens androgen receptor coactivator ARA55 mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF129530 AF129530 Homo sapiens chromosome 10 F-box protein Fbw1A (FBW1A) mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF144638 AF144638 Homo sapiens sphingosine-1-phosphate lyase (SPL) mRNA, complete cds.
Human beta-TrCP1	SEQ ID 8	gb AF164678 AF164678 Homo sapiens FLASH homolog RIP25 (RIP25) mRNA, complete

			cds.
Human beta-TrCP1	SEQ ID 8	gb AF167160 AF167160 Homo sapiens protein inhibitor of activated STAT-1 (PIAS1) mRNA, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb AF191298 AF191298 Homo sapiens vacuolar sorting protein 35 (VPS35) mRNA, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb AF250238 AF250238 Homo sapiens macrophage ABC transporter (ABCA7) mRNA, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb AJ010089 HSA010089 Homo sapiens mRNA for GANP protein.	
Human beta-TrCP1	SEQ ID 8	gb AJ242910 HSA242910 Homo sapiens mRNA for N-Acetylglucosamine kinase.	
Human beta-TrCP1	SEQ ID 8	gb AL035413 HS657E11 Human DNA sequence from clone RP4-657E11 on chromosome 1p35.1-36.23	
Human beta-TrCP1	SEQ ID 8	gb AL110226 HSM800883 Homo sapiens mRNA; cDNA DKFZp434H204 (from clone DKFZp434H204).	
Human beta-TrCP1	SEQ ID 8	gb AL137497 HSM802227 Homo sapiens mRNA; cDNA DKFZp761C241 (from clone DKFZp761C241).	
Human beta-TrCP1	SEQ ID 8	gb AL157477 HSM802464 Homo sapiens mRNA; cDNA DKFZp761E212 (from clone DKFZp761E212).	
Human beta-TrCP1	SEQ ID 8	gb AQ571615 AQ571615 HS_5380_B2_C03_SP0E RPCI-11 Human Male BAC Library Homo sapiens enomic clone Plate=956 Col=6 Row=F, DNA sequence.	
Human beta-TrCP1	SEQ ID 8	gb D17032 HUMD3D08M5 Human HepG2 partial cDNA, clone hmd3d08m5.	
Human beta-TrCP1	SEQ ID 8	gb D78360 HUMPP27KRS Homo sapiens mRNA for protein phosphatase 2A 74 kDa regulatory subunit (delta or B'' subunit), complete cds.	
Human beta-TrCP1	SEQ ID 8	gb D84295 D84295 Human mRNA for possible protein TPRDII, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb D87073 D87073 Human mRNA for KIAA0236 gene, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb G24929 G24929 human STS EST204289, sequence tagged site.	
Human beta-TrCP1	SEQ ID 8	gb J03210 HUMCN4GEL Human collagenase type IV mRNA, 3' end.	
Human beta-TrCP1	SEQ ID 8	gb K02581 HUMTK Human thymidine kinase mRNA, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb L32602 RATOTX1X Rattus norvegicus OTX1 mRNA, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb M55971 HPCNS2PA Human hepatitis virus C NS2 protein, partial cds.	
Human beta-TrCP1	SEQ ID 8	gb M67483 HPCCGAA Hepatitis C virus, complete genome.	
Human beta-TrCP1	SEQ ID 8	gb U33760 HSU33760 Human cyclin A/CDK2-associated p19 (Sklp1) mRNA, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb U45328 HSU45328 Human ubiquitin-conjugating enzyme (UBE2L) mRNA, complete cds.	
Human beta-TrCP1	SEQ ID 8	gb U51166 HSU51166 Human G/T mismatch-specific thymine DNA glycosylase mRNA, complete cds.	

human Rac1	SEQ ID 10	gb AC025580 AC025580 Homo sapiens chromosome 15 clone RP11-519G16 map 15q21, WORKING DRAFT SEQUENCE, 24 unordered pieces.
human Rac1	SEQ ID 10	gb AC026634 AC026634 Homo sapiens chromosome 18 clone RP11-639E23 map 18, WORKING DRAFT SEQUENCE, 23 unordered pieces.
human Rac1	SEQ ID 10	gb AC027493 AC027493 Homo sapiens chromosome 3 clone RP11-585F20 map 3, WORKING DRAFT SEQUENCE, 13 unordered pieces.
human Rac1	SEQ ID 10	gb AC031989 AC031989 Homo sapiens chromosome 6 clone RP11-183F17 map 6, WORKING DRAFT SEQUENCE, 20 unordered pieces.
human Rac1	SEQ ID 10	gb AC032022 AC032022 Homo sapiens chromosome 2 clone RP11-321C18 map 2, WORKING DRAFT SEQUENCE, 31 unordered pieces.
human Rac1	SEQ ID 10	gb AC069391 AC069391 Homo sapiens chromosome 7 clone RP11-462D19, WORKING DRAFT SEQUENCE, 23 unordered pieces.
human Rac1	SEQ ID 10	gb AF008591 AF008591 Homo sapiens Rac3 (RAC3) mRNA, complete cds.
human Rac1	SEQ ID 10	gb AF118838 AF118838 Homo sapiens citrin (SLC25A13) mRNA, complete cds.
human Rac1	SEQ ID 10	gb AF159442 AF159442 Homo sapiens phospholipid scramblase 3 mRNA, complete cds.
human Rac1	SEQ ID 10	gb AF283769 AF283769 Homo sapiens clone TCBA00758 mRNA sequence.
human Rac1	SEQ ID 10	gb AJ133269 HSA133269 Homo sapiens caveolin-1/-2 locus, Contig1, D7S522, genes CAV2 (exons 1, 2a, and 2b), CAV1 (exons 1 and 2).
human Rac1	SEQ ID 10	gb AL031775 HS30M3 Human DNA sequence from clone 30M3 on chromosome 6p22-22.3. Contains three novel genes, one similar to C. elegans Y63D3A.4 and one similar to (predicted) plant, worm, yeast and archaea bacterial genes, and the first exon of the KIAA0319 gene. Contains ESTs, GSSs and putative CpG islands, complete sequence.
human Rac1	SEQ ID 10	gb AL035413 HS657E11 Human DNA sequence from clone RP4-657E11 on chromosome 1p35-36.23 Contains 3' part of the CAPZB (capping protein (actin filament) muscle Z-line, beta) gene, genes for aldo-keto reductase family 7 (aflatoxin aldehyde reductase) mgbars A2 (AKR7A2) and A3 (AKR7A3), a novel gene similar to acidic ribosomal protein PO, the gene for KIAA0090 protein, ESTs, STSs, GSSs and CpG islands, complete sequence.
human Rac1	SEQ ID 10	gb AL110179 HSM800827 Homo sapiens mRNA; cDNA DKFZp564D0472 (from clone DKFZp564D0472).
human Rac1	SEQ ID 10	gb AL121898 HSA430K20 Human DNA sequence from clone RP11-430K20 on chromosome 20. Contains GSSs and a CpG island, complete sequence.
human Rac1	SEQ ID 10	gb AL139133 AL139133 Homo sapiens chromosome 1 clone RP11-185C19 map q31-31.3
human Rac1	SEQ ID 10	gb AL139188 AL139188 Human DNA sequence from clone RP11-90M5 on chromosome 13q121-12.3, complete sequence.

human Rac1	SEQ ID 10	gb AL139812 AL139812 Human DNA sequence from clone RP1-19N1 on chromosome Xq21.33-22.3, complete sequence.
human Rac1	SEQ ID 10	gb AL157477 HSM802464 Homo sapiens mRNA; cDNA DKFZp761E212 (from clone DKFZp761E212).
human Rac1	SEQ ID 10	gb AL162272 AL162272 Homo sapiens chromosome 10 clone RP11-145E8
human Rac1	SEQ ID 10	gb AL353800 AL353800 Homo sapiens chromosome 6 clone RP3-442A17
human Rac1	SEQ ID 10	gb AL359611 HSM802730 Homo sapiens mRNA; cDNA DKFZp762L1710 (from clone DKFZp762L1710).
human Rac1	SEQ ID 10	gb AP000356 AP000356 Homo sapiens genomic DNA, chromosome 22q11.2, clone KB1995A5.
human Rac1	SEQ ID 10	gb AP001385 AP001385 Homo sapiens chromosome 11 clone RP11-669D23 map 11q13, WORKING DRAFT SEQUENCE, in unordered pieces.
human Rac1	SEQ ID 10	gb AQ533697 AQ533697 RPCI-11-384A12.TV RPCI-11 Homo sapiens genomic clone PCI-11-384A12, DNA sequence.
human Rac1	SEQ ID 10	gb D50911 D50911 Homo sapiens mRNA for KIAA0121 protein, partial cds.
human Rac1	SEQ ID 10	gb L25080 HUMRH0AA Homo sapiens GTP-binding protein (rhoA) mRNA, complete cds.
human Rac1	SEQ ID 10	gb M29870 HUMRACA Human ras-related C3 botulinum toxin substrate (rac) mRNA, complete cds.
human Rac1	SEQ ID 10	gb M33519 HUMBAT3A Human HLA-B-associated transcript 3 (BAT3) mRNA, complete cds.
human Rac1	SEQ ID 10	gb M57298 HUMGPG25K Human GTP-binding protein G25K mRNA, complete cds.
human Rac1	SEQ ID 10	gb M67463 HPCCGAA Hepatitis C virus, complete genome.
human Rac1	SEQ ID 10	gb U55017 HSU55017 Human transketolase (TKT) mRNA, complete cds.
human Rac1	SEQ ID 10	gb U62317 HUB384D8 Homo sapiens chromosome 22q13 BAC clone CIT987SK-384D8 complete sequence.
human Rac1	SEQ ID 10	gb U91327 HSU91327 Human chromosome 12p15 BAC clone CIT987SK-99D8 complete sequence.
human Rac1	SEQ ID 10	gb X52022 HSCOLLV13 H.sapiens RNA for type VI collagen alpha3 chain.
human Rac1	SEQ ID 10	gb X54486 HSC1INH1B Human gene for C1-inhibitor.
human Rac1	SEQ ID 10	gb X61587 HSRHOG H.sapiens rhoG mRNA for GTPase.
human Rac1	SEQ ID 10	gb X86691 HSM12218 H.sapiens mRNA for 218kD Mi-2 protein.
Human uracil DNA glycosylase	SEQ ID 12	gb AB011089 AB011089 Homo sapiens mRNA for KIAA0517 protein, partial cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AB011121 AB011121 Homo sapiens mRNA for KIAA0549 protein, partial cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AB020638 AB020638 Homo sapiens mRNA for KIAA0831 protein, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AB037839 AB037839 Homo sapiens mRNA for KIAA1418 protein, partial cds.

Human uracil DNA glycosylase	SEQ ID 12	gb AB037856 AB037856 Homo sapiens mRNA for KIAA1435 protein, partial cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AC002546 AC002546 Homo sapiens chromosome 17, clone 193h18, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AC003108 HUAC003108 Human Chromosome 16 BAC clone CIT987SK-327024, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AC003681 AC003681 Human PAC clone RP3-394A18 from 22q12-qter, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AC004850 AC004850 Homo sapiens PAC clone RP4-665C4 from 7p14-p13, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AC005820 AC005820 Homo sapiens chromosome Y, clone hCIT.494_G_17, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AC006530 AC006530 Homo sapiens chromosome 14 clone BAC 316E14 map 14q24.3, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AC007386 AC007386 Homo sapiens BAC clone RP11-359K10 from 2, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AC007779 AC007779 Homo sapiens chromosome 9 clone RP11-483H20 map 9.
Human uracil DNA glycosylase	SEQ ID 12	gb AC008570 AC008570 Homo sapiens chromosome 5 clone CTC-549A4
Human uracil DNA glycosylase	SEQ ID 12	gb AC011236 AC011236 Homo sapiens chromosome 2 clone RP11-312D1
Human uracil DNA glycosylase	SEQ ID 12	gb AC012014 AC012014 Homo sapiens chromosome 3 clone RP11-255N4
Human uracil DNA glycosylase	SEQ ID 12	gb AC012278 AC012278 Homo sapiens chromosome 11 clone RP11-35D12 map 11
Human uracil DNA glycosylase	SEQ ID 12	gb AC012342 AC012342 Homo sapiens chromosome 3 clone RP11-436F13 map 3
Human uracil DNA glycosylase	SEQ ID 12	gb AC016750 AC016750 Homo sapiens chromosome 11 clone RP11-504G3
Human uracil DNA glycosylase	SEQ ID 12	gb AC019070 AC019070 Homo sapiens chromosome 2 clone RP11-264M11
Human uracil DNA glycosylase	SEQ ID 12	gb AC022816 AC022816 Homo sapiens chromosome 17 clone RP11-626C5 map 17
Human uracil DNA glycosylase	SEQ ID 12	gb AC023785 AC023785 Homo sapiens chromosome 6 clone RP11-507C10
Human uracil DNA glycosylase	SEQ ID 12	gb AC024242 AC024242 Homo sapiens chromosome 8 clone RP11-513D5
Human uracil DNA glycosylase	SEQ ID 12	gb AC025192 AC025192 Homo sapiens chromosome 8 clone RP11-509E2 map 8
Human uracil DNA glycosylase	SEQ ID 12	gb AC025817 AC025817 Homo sapiens chromosome 3 clone RP11-143O1
Human uracil DNA glycosylase	SEQ ID 12	gb AC031989 AC031989 Homo sapiens chromosome 6 clone RP11-183F17 map 6
Human uracil DNA glycosylase	SEQ ID 12	gb AF155120 AF155120 Homo sapiens ubiquitin-conjugating enzyme variant Kua (UBE2V) mRNA, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AF156857 AF156857 Homo sapiens actin-binding protein (lPP) mRNA, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AF178980 AF178980 Homo sapiens D-prohibitin mRNA, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AF191298 AF191298 Homo sapiens vacuolar sorting protein 35 (VPS35) mRNA, complete cds.

Human uracil DNA glycosylase	SEQ ID 12	gb AF224669 AF213884 S2 Homo sapiens mannosidase, beta A, lysosomal (MANBA) gene, and ubiquitin-conjugating enzyme E2D 3 (UBE2D3) genes, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb AJ133269 HSA133269 Homo sapiens caveolin-1/-2 locus, Contig1, D7S522, genes CAV2 (exons 1, 2a, and 2b), CAV1 (exons 1 and 2).
Human uracil DNA glycosylase	SEQ ID 12	gb AL009443 HSP11B05 H.sapiens flow-sorted chromosome 1 HindIII fragment, SC1pE11B05, sequence tagged site.
Human uracil DNA glycosylase	SEQ ID 12	gb AL031577 HS391O22 Human DNA sequence from clone 391O22 on chromosome 6p21.2-21.31. Contains pseudogenes similar to ribosomal proteins L44 and L30, a pseudogene similar to interferon-inducible protein 1-8U, ESTs, GSSs, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AL109797 HS1172N10 Human DNA sequence from clone RP5-1172N10 on chromosome Xp11.3-11.4, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AL118511 HSDJ858B6 Homo sapiens chromosome 1 clone RP5-858B6 map q423-43
Human uracil DNA glycosylase	SEQ ID 12	gb AL139133 AL139133 Homo sapiens chromosome 1 clone RP11-185C19 map q31-31.3
Human uracil DNA glycosylase	SEQ ID 12	gb AL139188 AL139188 Human DNA sequence from clone RP11-90M5 on chromosome 13q12-12.3, complete sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb AL157477 HSM802464 Homo sapiens mRNA; cDNA DKFZp761E212 (from clone DKFZp761E212).
Human uracil DNA glycosylase	SEQ ID 12	gb AL157894 AL157894 Homo sapiens chromosome 10 clone RP11-325E17.
Human uracil DNA glycosylase	SEQ ID 12	gb AL355885 CNS05TCW Homo sapiens clone R-43A022
Human uracil DNA glycosylase	SEQ ID 12	gb AL359611 HSM802730 Homo sapiens mRNA; cDNA DKFZp762L1710 (from clone DKFZp762L1710).
Human uracil DNA glycosylase	SEQ ID 12	gb AL365208 AL365208 Homo sapiens chromosome 1 clone RP4-706A17
Human uracil DNA glycosylase	SEQ ID 12	gb AP000356 AP000356 Homo sapiens genomic DNA, chromosome 22q11.2, clone KB1995A5.
Human uracil DNA glycosylase	SEQ ID 12	gb AQ771795 AQ771795 HS_5410_B1_E01_T7A RPCL-11 Human Male BAC Library Homo sapiens genomic clone Plate=986 Col=1 Row=J, DNA sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb B39132 B39132 HS-1049-A1-E12-MR.abi CIT Human Genomic Sperm Library C Homo sapiens genomic clone Plate=CT 771 Col=23 Row=I, DNA sequence.
Human uracil DNA glycosylase	SEQ ID 12	gb D31889 HUMORFKG1R Human mRNA for KIAA0072 gene, partial cds.
Human uracil DNA glycosylase	SEQ ID 12	gb G24929 G24929 human STS EST204289, sequence tagged site.
Human uracil DNA glycosylase	SEQ ID 12	gb J05249 HUMREPA Human replication protein A 32-kDa subunit mRNA, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb J16279 HUMMIC2A Human MIC2 mRNA, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb J16447 HUMDHPRA Human dihydropteridine reductase (hDHPR) mRNA, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb J1724 HUMPTPBX Human phosphotyrosyl-protein phosphatase (PTP-1B) mRNA,

Human uracil DNA glycosylase	SEQ ID 12	complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb S77127 S77127 Homo sapiens manganese superoxide dismutase gene, complete cds.
		gb U43195 HSU43195 Human Rho-associated, coiled-coil containing protein kinase p160ROCK mRNA, complete cds.
Human uracil DNA glycosylase	SEQ ID 12	gb Z83844 HS37E16 Human DNA sequence from clone RP1-37E16 on chromosome 22
Human b2 adrenergic receptor	SEQ ID 14	gb AB002360 AB002360 Human mRNA for KIAA0362 gene, partial cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AB002370 AB002370 Human mRNA for KIAA0372 gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AB007890 AB007890 Homo sapiens mRNA for KIAA0430 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AB018306 AB018306 Homo sapiens mRNA for KIAA0763 protein, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AB028956 AB028956 Homo sapiens mRNA for KIAA1033 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AB029290 AB029290 Homo sapiens mRNA for actin binding protein ABP620, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AB033076 AB033076 Homo sapiens mRNA for KIAA1250 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AC002366 AC002366 Human Xp22 BAC CT-285115 (from CalTech/Research Genetics) , PAC RPC11-27C22 (from Roswell Park Cancer Center), and Cosmid U35B5 (from Lawrence Livermore), complete sequence.
Human b2 adrenergic receptor	SEQ ID 14	gb AF001893 BETA2 Human MEN1 region clone epsilon/beta mRNA, 3' fragment.
Human b2 adrenergic receptor	SEQ ID 14	gb AF024631 AF024631 Homo sapiens ANG2 (ANG2) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AF061738 AF061738 Homo sapiens leucine aminopeptidase mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AF098799 AF098799 Homo sapiens RanBP7/importin 7 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AF124490 AF124490 Homo sapiens ARF GTPase-activating protein GIT1 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AF128862 AF128862 Cloning vector pHIND2.2, complete sequence.
Human b2 adrenergic receptor	SEQ ID 14	gb AF151815 AF151815 Homo sapiens CGI-57 protein mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb AL035413 HS657E11 Human DNA sequence from clone RP4-657E11 on chromosome 1p35.1-36.23
Human b2 adrenergic receptor	SEQ ID 14	gb AL121820 CNS01DSM Human chromosome 14 DNA, BAC C-2325P2 of library CalTech-D from chromosome 14 of Homo sapiens (Human), complete sequence.
Human b2 adrenergic receptor	SEQ ID 14	gb AL157419 HSM802422 Homo sapiens mRNA; cDNA DKFZp434P031 (from clone DKFZp434P031).
Human b2 adrenergic receptor	SEQ ID 14	gb AQ175201 AQ175201 HS_3212_B2_F05_T7 CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=3212 Col=10 Row=L, DNA sequence.
Human b2 adrenergic receptor	SEQ ID 14	gb D28476 HUMKG1C Human mRNA for KIAA0045 gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb L32602 RATOTX1X Rattus norvegicus OTX1 mRNA, complete cds.

Human b2 adrenergic receptor	SEQ ID 14	gb L76703 HUMB56EA Homo sapiens protein phosphatase 2A B56-epsilon (PP2A) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb M60119 HUMEP2AA Homo sapiens HIV-EP2/Schnurri-2 gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb S66431 S66431 RBP2=retinoblastoma binding protein 2 [human, Nalm-6 pre-B cell leukemia, mRNA, 6455 nt].
Human b2 adrenergic receptor	SEQ ID 14	gb U06863 HSU06863 Human follistatin-related protein precursor mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb U28964 HSU28964 Homo sapiens 14-3-3 protein mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb U56998 HSU56998 Human putative serine/threonine protein kinase PRK (prk) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb U59289 HSU59289 Human H-cadherin mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb U65928 HSU65928 Human Jun activation domain binding protein mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb U89867 HSU89867 Human nuclear matrix protein 55 (nmt55) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 14	gb X53416 HSABP280 Human mRNA for actin-binding protein (filamin) (ABP-280).
Human b2 adrenergic receptor	SEQ ID 16	gb AB011164 AB011164 Homo sapiens mRNA for KIAA0592 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AB019691 AB019691 Homo sapiens mRNA for Centrosome- and Golgi-localized PKN-associated protein (CG-NAP), complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AB032252 AB032252 Homo sapiens BAZ1A mRNA for bromodomain adjacent to zinc finger domain 1A, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AB033094 AB033094 Homo sapiens mRNA for KIAA1268 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AC009336 AC009336 Homo sapiens chromosome 2, clone RP11-387A1, complete sequence.
Human b2 adrenergic receptor	SEQ ID 16	gb AF001893 BETA2 Human MEN1 region clone epsilon/beta mRNA, 3' fragment.
Human b2 adrenergic receptor	SEQ ID 16	gb AF021935 AF021935 Rattus norvegicus myotonic dystrophy kinase-related Cdc42-binding kinase (MRCK) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AF098638 AF098638 Homo sapiens rabaptin-4 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AF117107 AF117107 Homo sapiens IGF-II mRNA-binding protein 2 (IMP-2) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AF174498 AF174498 Homo sapiens GR AF-1 specific protein phosphatase mRNA, partial cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AF191298 AF191298 Homo sapiens vacuolar sorting protein 35 (VPS35) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb AJ010089 HSA010089 Homo sapiens mRNA for GANP protein.
Human b2 adrenergic receptor	SEQ ID 16	gb AJ278112 HSA278112 Homo sapiens mRNA for putative cell cycle control protein (SDP35 gene).

Human b2 adrenergic receptor	SEQ ID 16	gb AK000046 AK000046 Homo sapiens cDNA FLJ20039 fis, clone COL00364.
Human b2 adrenergic receptor	SEQ ID 16	gb AK001434 AK001434 Homo sapiens cDNA FLJ10572 fis, clone NT2RP2003125, weakly similar to RING CANAL PROTEIN.
Human b2 adrenergic receptor	SEQ ID 16	gb AL078633 HS1005F21 Human DNA sequence from clone RP5-1005F21 on chromosome 20, complete sequence.
Human b2 adrenergic receptor	SEQ ID 16	gb AL117187 CNS01DRD Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-725G5 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
Human b2 adrenergic receptor	SEQ ID 16	gb AL352979 CNS05TBZ Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-368P15 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
Human b2 adrenergic receptor	SEQ ID 16	gb AP000349 AP000349 Homo sapiens genomic DNA, chromosome 22q11.2, clone KB1839H6.
Human b2 adrenergic receptor	SEQ ID 16	gb AQ345186 AQ345186 RPCI11-123C5.TV RPCI-11 Homo sapiens genomic clone RPCI-11-123C5, DNA sequence.
Human b2 adrenergic receptor	SEQ ID 16	gb AQ376051 AQ376051 RPCI11-150L20.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-150L20, DNA sequence.
Human b2 adrenergic receptor	SEQ ID 16	gb AQ787273 AQ787273 HS_5565_B1_H03_T7A RPCI-11 Human Male BAC Library Homo sapiens genomic clone Plate=1141 Col=5 Row=P, DNA sequence.
Human b2 adrenergic receptor	SEQ ID 16	gb D38047 HUMPSP31 Human mRNA for 26S proteasome subunit p31, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb J01415 HUMMTG Human mitochondrion, complete genome.
Human b2 adrenergic receptor	SEQ ID 16	gb J02959 HUMMLKHA4 Human leukotriene A-4 hydrolase mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb J03077 HUMGLBA Human co-beta glucosidase (proactivator) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb J04177 HUMCA1XIA Human alpha-1 type XI collagen (COL11A1) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb L78810 HUMYWXD703 Homo sapiens ADP/ATP carrier protein (ANT-2) gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb M12529 HUMAPOE Human apolipoprotein E mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb M32221 HUMSAPABCD Human saposin proteins A-D mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb M86667 HUMNAP H.sapiens NAP (nucleosome assembly protein) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb M96803 HUMSPTBN1A Human general beta-spectrin (SPTBN1) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb U06863 HSU06863 Human follistatin-related protein precursor mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb U12596 HSU12596 Human tumor necrosis factor type 1 receptor associated protein (TRAP2) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb U15641 HSU15641 Human transcription factor E2F-4 mRNA, complete cds.

Human b2 adrenergic receptor	SEQ ID 16	gb U22384 HSU22384 Human lysyl oxidase gene, partial cds.
Human b2 adrenergic receptor	SEQ ID 16	gb U29343 HSU29343 Homo sapiens hyaluronan receptor (RHAMM) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb U41766 HSU41766 Human metalloprotease/disintegrin/cysteine-rich protein precursor (MDC9) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb U50532 HSU50532 Human BRCA2 region, mRNA sequence CG005.
Human b2 adrenergic receptor	SEQ ID 16	gb U67280 HSU67280 Homo sapiens calumenin mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 16	gb X53416 HSABP280 Human mRNA for actin-binding protein (filamin) (ABP-280).
Human b2 adrenergic receptor	SEQ ID 16	gb Y11997 HSY11997 H.sapiens mRNA for A-kinase anchoring protein AKAP95.
Human b2 adrenergic receptor	SEQ ID 18	gb AB002319 AB002319 Human mRNA for KIAA0321 gene, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB002340 AB002340 Homo sapiens mRNA for KIAA0342 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB002360 AB002360 Human mRNA for KIAA0362 gene, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB002370 AB002370 Human mRNA for KIAA0372 gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB006629 AB006629 Homo sapiens mRNA for KIAA0291 gene, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB006757 AB006757 Homo sapiens mRNA for PCDH7 (BH-Pcdh)c, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB007890 AB007890 Homo sapiens mRNA for KIAA0430 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB018271 AB018271 Homo sapiens mRNA for KIAA0728 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB018313 AB018313 Homo sapiens mRNA for KIAA0770 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB023224 AB023224 Homo sapiens mRNA for KIAA1007 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB024334 AB024334 Homo sapiens mRNA for 14-3-3gamma, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB028956 AB028956 Homo sapiens mRNA for KIAA1033 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB028981 AB028981 Homo sapiens mRNA for KIAA1058 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB028990 AB028990 Homo sapiens mRNA for KIAA1067 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB029290 AB029290 Homo sapiens mRNA for actin binding protein ABP620, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB032253 AB032253 Homo sapiens BAZ1B mRNA for bromodomain adjacent to zinc finger domain 1B, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB035898 AB035898 Homo sapiens hklp2 mRNA for kinesin-like protein 2, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AB037850 AB037850 Homo sapiens mRNA for KIAA1429 protein, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AC010553 AC010553 Homo sapiens chromosome 16 clone RP11-59D8, complete sequence.
Human b2 adrenergic receptor	SEQ ID 18	gb AF006751 AF006751 Homo sapiens ES/130 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF019225 AF019225 Homo sapiens apolipoprotein L mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF039023 AF039023 Homo sapiens Ran-GTP binding protein mRNA, partial cds.

Human b2 adrenergic receptor	SEQ ID 18	gb AF049527 AF049527 Homo sapiens alpha-adaptin C mRNA, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF054589 AF054589 Homo sapiens HIC protein isoform p40 and HIC protein isoform p32 mRNAs, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF061555 AF061555 Mus musculus ubiquitin-protein ligase E3-alpha (Ubr1) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF061738 AF061738 Homo sapiens leucine aminopeptidase mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF084457 AF084457 Homo sapiens beta-cop homolog mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF093543 AF093543 Homo sapiens transforming acidic coiled-coil containing protein 3 (TACC3) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF094519 AF094519 Mus musculus diaphanous-related formin (Dia2) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF124490 AF124490 Homo sapiens ARF GTPase-activating protein GIT1 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF128625 AF128625 Homo sapiens CDC42-binding protein kinase beta (CDC42BPB) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF151857 AF151857 Homo sapiens CGI-99 protein mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF156102 AF156102 Homo sapiens ELL complex EAP30 subunit mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF198092 AF198092 Mus musculus RP42 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF201390 AF201390 Mus musculus p300 transcriptional cofactor JMY mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF216185 AF216185 Homo sapiens tissue-type brain Ellis-van Creveld syndrome protein (EVC) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF217796 AF217796 Homo sapiens SCG10 like-protein, helicase-like protein NHL, M68, and ADP-ribosylation factor related protein 1 (ARFRP1) genes, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AF242319 AF242319 Mus musculus unknown mRNA.
Human b2 adrenergic receptor	SEQ ID 18	gb AJ010089 HSA010089 Homo sapiens mRNA for GANP protein.
Human b2 adrenergic receptor	SEQ ID 18	gb AK000393 AK000393 Homo sapiens cDNA FLJ20386 fis, clone KAlA4184.
Human b2 adrenergic receptor	SEQ ID 18	gb AL050101 HSM800186 Homo sapiens mRNA; cDNA DKFZp586E1519 (from clone DKFZp586E1519); partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb AL133245 CNS01DUI BAC sequence from the SPG4 candidate region at 2p21-2p22 BAC 854M03 of RPCI-11 library from chromosome 2 of Homo sapiens (Human), complete sequence.
Human b2 adrenergic receptor	SEQ ID 18	gb AL360275 IR1672402 Homo sapiens mRNA full length insert cDNA clone EUROIMAGE 1672402.

Human b2 adrenergic receptor	SEQ ID 18	gb AP000124 AP000124 Homo sapiens genomic DNA of 21q22.1, GART and AML related, SLC5A3-4A4 region, segment 7/8, complete sequence.
Human b2 adrenergic receptor	SEQ ID 18	gb AQ895869 AQ895869 HS_5524_B1_A06_T7A RPCI-11 Human Male BAC Library Homo sapiens genomic clone Plate=9292 Col=11 Row=B, DNA sequence.
Human b2 adrenergic receptor	SEQ ID 18	gb AR060756 AR060756 Sequence 2 from patent US 5840866.
Human b2 adrenergic receptor	SEQ ID 18	gb D26361 HUMORFW Human mRNA for KIAA0042 gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb D28476 HUMKG1C Human mRNA for KIAA0045 gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb D83781 D83781 Human mRNA for KIAA0197 gene, partial cds.
Human b2 adrenergic receptor	SEQ ID 18	gb J05032 HUMASP Human aspartyl-tRNA synthetase alpha-2 subunit mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb L20422 HUM1433ACT Human 14-3-3n protein mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb L32602 RATOTX1X Rattus norvegicus OTX1 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb L33075 HUMIQGA Homo sapiens ras GTPase-activating-like protein (IQGAP1) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb L39793 HUMNTF9 Homo sapiens nuclear factor p97 (NTF97) gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb L42572 HUMP8789R Homo sapiens p87/89 gene, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb L76703 HUMB56EA Homo sapiens protein phosphatase 2A B56-epsilon (PP2A) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb M12529 HUMAPOE Human apolipoprotein E mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb M95178 HUMACTN1A Human non-muscle alpha-actinin mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb M96803 HUMSPTBN1A Human general beta-spectrin (SPTBN1) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U06863 HSU06863 Human follistatin-related protein precursor mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U19721 HSU19721 Human peroxisomal targeting signal receptor 1 (PXR1) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U28964 HSU28964 Homo sapiens 14-3-3 protein mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U29343 HSU29343 Homo sapiens hyaluronan receptor (RHAMM) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U35113 HSU35113 Human metastasis-associated mita1 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U42390 HSU42390 Homo sapiens Trio mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U53204 HSU53204 Human plectin (PLEC1) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U54778 HSU54778 Human 14-3-3 epsilon mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U59289 HSU59289 Human H-cadherin mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U69139 HSU69139 Human zyglin mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U72761 HSU72761 Human karyopherin beta 3 mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 18	gb U83867 HSU83867 Human alpha II spectrin mRNA, complete cds.

Human b2 adrenergic receptor	SEQ ID 20	gb U46571 HSU46571 Human tetratricopeptide repeat protein (tpr2) mRNA, complete cds.
Human b2 adrenergic receptor	SEQ ID 20	gb U47077 HSU47077 Homo sapiens DNA-dependent protein kinase catalytic subunit (DNA-PKcs) mRNA, complete cds.
hSHP2_FL	SEQ ID 22	gb AL133367 CNS01DUS Human chromosome 14 DNA sequence BAC R-600F24 of library RPC1-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
	SEQ ID 22	gb AQ018017 AQ018017 CIT-HSP-2307B6.TR CIT-HSP Homo sapiens genomic clone 2307B6, DNA sequence.
	SEQ ID 22	gb U61843 HSU61843 Human discs large protein P-dlg mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AB000280 AB000280 Rattus norvegicus mRNA for peptide/histidine transporter, complete cds.
Human OBRGRP	SEQ ID 24	gb AB004788 AB004788 Homo sapiens mRNA for BNIP3L, complete cds.
Human OBRGRP	SEQ ID 24	gb AB011169 AB011169 Homo sapiens mRNA for KIAA0597 protein, partial cds.
Human OBRGRP	SEQ ID 24	gb AB015355 AB015355 Homo sapiens NRAMP2 gene for natural resistance-associated macrophage protein 2, complete cds.
Human OBRGRP	SEQ ID 24	gb AB015856 AB015856 Homo sapiens mRNA for ATF6, complete cds.
Human OBRGRP	SEQ ID 24	gb AB018010 AB018010 Homo sapiens mRNA for 4F2 heavy chain, complete cds.
Human OBRGRP	SEQ ID 24	gb AB018310 AB018310 Homo sapiens mRNA for KIAA0767 protein, partial cds.
Human OBRGRP	SEQ ID 24	gb AB019002 AB019002 Homo sapiens MRP5 mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AB020980 AB020980 Homo sapiens mRNA for putative membrane protein, complete cds.
Human OBRGRP	SEQ ID 24	gb AB032995 AB032995 Homo sapiens mRNA for KIAA1169 protein, partial cds.
Human OBRGRP	SEQ ID 24	gb AB033091 AB033091 Homo sapiens mRNA for KIAA1265 protein, partial cds.
Human OBRGRP	SEQ ID 24	gb AC002398 AC002398 Human DNA from chromosome 19-specific cosmid F25965, genomic sequence, complete sequence.
Human OBRGRP	SEQ ID 24	gb AC003101 AC003101 Homo sapiens chromosome 17, clone HRP41C23, complete sequence.
Human OBRGRP	SEQ ID 24	gb AC007099 AC007099 Homo sapiens BAC clone RP11-445A14 from 2, complete sequence.
Human OBRGRP	SEQ ID 24	gb AC007279 AC007279 Homo sapiens clone NH0309N08, complete sequence.
Human OBRGRP	SEQ ID 24	gb AC009505 AC009505 Homo sapiens clone RP11-526D2, complete sequence.
Human OBRGRP	SEQ ID 24	gb AF002697 AF002697 Homo sapiens E1B 19K/Bcl-2-binding protein Nip3 mRNA, nuclear gene encoding mitochondrial protein, complete cds.
Human OBRGRP	SEQ ID 24	gb AF005039 AF005039 Homo sapiens secretory carrier membrane protein (SCAMP3) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF053755 AF053755 Homo sapiens bicarbonate transporter (BT) mRNA, complete cds.

Human OBRGRP	SEQ ID 24	gb AF069512 AF069512 Homo sapiens sodium bicarbonate cotransporter (NBC) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF081282 AF081282 Homo sapiens small membrane protein 1 (SMP1) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF097535 AF097535 Homo sapiens membrane protein CH1 (CH1) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF105365 AF105365 Homo sapiens K-Cl cotransporter KCC4 mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF117330 AF117330 Rattus norvegicus unknown mRNA.
Human OBRGRP	SEQ ID 24	gb AF126799 AF126799 Homo sapiens delta-6 fatty acid desaturase mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF147742 AF147742 Homo sapiens myeloid cell differentiation protein (MCL1) gene, promoter and complete cds.
Human OBRGRP	SEQ ID 24	gb AF151799 AF151799 Homo sapiens CGI-40 protein mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF161424 AF161424 Homo sapiens HSPC306 mRNA, partial cds.
Human OBRGRP	SEQ ID 24	gb AF161446 AF161446 Homo sapiens HSPC328 mRNA, partial cds.
Human OBRGRP	SEQ ID 24	gb AF167706 AF167706 Homo sapiens cysteine-rich repeat-containing protein S52 precursor, mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF198097 AF198097 Homo sapiens chromosome Xp11.23 cosmid B167, E1017, and L2460, complete sequence.
Human OBRGRP	SEQ ID 24	gb AF209704 AF209704 Homo sapiens glycolipid transfer protein mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF273024 AF273024 Rattus norvegicus amino acid system A transporter mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AF285167 AF285167 Homo sapiens ATP-binding cassette transporter 1 (ABCA1) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb AJ011863 HSA011863 Homo sapiens mRNA for homeobox protein LSX.
Human OBRGRP	SEQ ID 24	gb AJ243936 HSA243936 Homo sapiens mRNA for G16 protein (G16 gene located in the class III region of the major histocompatibility complex).
Human OBRGRP	SEQ ID 24	gb AK000060 AK000060 Homo sapiens cDNA FLJ20053 fis, clone COL00809.
Human OBRGRP	SEQ ID 24	gb AK000331 AK000331 Homo sapiens cDNA FLJ20324 fis, clone HEP09841, highly similar to AB007931 Homo sapiens mRNA for KIAA0462 protein.
Human OBRGRP	SEQ ID 24	gb AK000630 AK000630 Homo sapiens cDNA FLJ20623 fis, clone KAT04793.
Human OBRGRP	SEQ ID 24	gb AK001571 AK001571 Homo sapiens cDNA FLJ10709 fis, clone NT2RP3000869.
Human OBRGRP	SEQ ID 24	gb AL110179 HSM800827 Homo sapiens mRNA; cDNA DKFZp564D0472 (from clone DKFZp564D0472).
Human OBRGRP	SEQ ID 24	gb AL360136 IR2176457 Homo sapiens mRNA full length insert cDNA clone EUROIMAGE 2176457.

Human OBRGRP	SEQ ID 24	gb AQ349348 AQ349348 RPC11-1139L5.TV RPC1-11 Homo sapiens genomic clone RPC1-11-139L5, DNA sequence.
Human OBRGRP	SEQ ID 24	gb B80296 B80296 CIT-HSP-2045D21.TR CIT-HSP Homo sapiens genomic clone 2045D21, DNA sequence.
Human OBRGRP	SEQ ID 24	gb D14582 HUMEP1 Human mRNA for epimorphin.
Human OBRGRP	SEQ ID 24	gb D30756 HUMORFKG11 Human mRNA for KIAA0049 gene, complete cds.
Human OBRGRP	SEQ ID 24	gb D38551 HUMORF005 Human mRNA for KIAA0078 gene, complete cds.
Human OBRGRP	SEQ ID 24	gb D50683 D50683 Homo sapiens mRNA for TGF-beta1R alpha, complete cds.
Human OBRGRP	SEQ ID 24	gb D87436 D87436 Human mRNA for KIAA0249 gene, complete cds.
Human OBRGRP	SEQ ID 24	gb D87742 D87742 Human mRNA for KIAA0268 gene, partial cds.
Human OBRGRP	SEQ ID 24	gb G24481 G24481 human STS WI-13464, sequence tagged site.
Human OBRGRP	SEQ ID 24	gb G24929 G24929 human STS EST204289, sequence tagged site.
Human OBRGRP	SEQ ID 24	gb J04027 HUMPMPCA Human plasma membrane Ca2+ pumping ATPase mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb L06133 HUMATPCU Human putative Cu++-transporting P-type ATPase mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb L21934 HUMACYLCOA Homo sapiens acyl-coenzyme A: cholesterol acyltransferase mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M16230 SUSMP1 Strongylocentrotus purpuratus spicule matrix protein SM37, partial cds; and spicule matrix protein SM50 precursor, gene, exon 1.
Human OBRGRP	SEQ ID 24	gb M16279 HUMMIC2A Human MIC2 mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M16965 HUMFP Human cerebellar degeneration-associated protein mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M23115 HUMHK2A Homo sapiens calcium-ATPase (HK2) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M29696 HUMIL7AA Human interleukin-7 receptor (IL-7) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M31627 HUMHXB1 Human X box binding protein-1 (XBP-1) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M55543 HUMGBP2 Human guanylate binding protein isoform II (GBP-2) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M64098 HUMHBP Human high density lipoprotein binding protein (HBP) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb M86737 HUMHMBP Human high mobility group box (SSRP1) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U00968 U00968 Human SREBP-1 mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U01874 HSU01874 Human me20m mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U02031 HSU02031 Human sterol regulatory element binding protein-2 mRNA, complete cds.

Human OBRGRP	SEQ ID 24	gb U17133 RNU17133 Rattus norvegicus zinc transporter ZnT-1 (ZnT-1) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U31903 HSU31903 Human CREB-RP (creb-rp) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U41060 HSU41060 Human breast cancer, estrogen regulated LIV-1 protein (LIV-1) mRNA, partial cds.
Human OBRGRP	SEQ ID 24	gb U41804 HSU41804 Human putative T1/ST2 receptor binding protein precursor mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U50939 HSU50939 Human amyloid precursor protein-binding protein 1 mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U51677 HSU51677 Human non-histone chromatin protein HMG1 (HMG1) gene, complete cds.
Human OBRGRP	SEQ ID 24	gb U55054 HSKCC Human K-Ci cotransporter (hKCC1) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U68063 HSU68063 Human transformer-2 beta (htra-2 beta) mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb U86755 HSU86755 Human TNF-alpha converting enzyme mRNA, complete cds.
Human OBRGRP	SEQ ID 24	gb X52425 HSL4R Human IL-4-R mRNA for the interleukin 4 receptor.
Human OBRGRP	SEQ ID 24	gb X53416 HSABP280 Human mRNA for actin-binding protein (filamin) (ABP-280).
Human OBRGRP	SEQ ID 24	gb X57398 HSPM5 Human mRNA for pM5 protein.
Human OBRGRP	SEQ ID 24	gb X98654 HSDRES9 H.sapiens mRNA for DRES9 protein.
Human OBRGRP	SEQ ID 24	gb Z83822 HS306D1 Human DNA sequence from PAC 306D1 on chromosome X contains ESTs.
Human OBRGRP	SEQ ID 26	gb AB002370 AB002370 Human mRNA for KIAA0372 gene, complete cds.
Human OBRGRP	SEQ ID 26	gb AB006651 AB006651 Homo sapiens EXLM1 mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AB008430 AB008430 Homo sapiens mRNA for CDEP, complete cds.
Human OBRGRP	SEQ ID 26	gb AB011472 AB011472 Homo sapiens mRNA for CDC23, complete cds.
Human OBRGRP	SEQ ID 26	gb AB012190 AB012190 Homo sapiens mRNA for Nedd8-activating enzyme hUba3, complete cds.
Human OBRGRP	SEQ ID 26	gb AB020718 AB020718 Homo sapiens mRNA for KIAA0911 protein, complete cds.
Human OBRGRP	SEQ ID 26	gb AB027196 AB027196 Homo sapiens mRNA for RIE2 sid2705, complete cds.
Human OBRGRP	SEQ ID 26	gb AB028956 AB028956 Homo sapiens mRNA for KIAA1033 protein, partial cds.
Human OBRGRP	SEQ ID 26	gb AB032966 AB032966 Homo sapiens mRNA for KIAA1140 protein, partial cds.
Human OBRGRP	SEQ ID 26	gb AB033034 AB033034 Homo sapiens mRNA for KIAA1208 protein, partial cds.
Human OBRGRP	SEQ ID 26	gb AF010404 AF010404 Homo sapiens ALR mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF021351 AF021351 Homo sapiens RNA polymerase III largest subunit (hRPC155) mRNA, complete cds.

Human OBRGRP	SEQ ID 26	gb AF024694 AF024694 Homo sapiens clone B3-3 zinc finger protein mRNA, partial cds.
Human OBRGRP	SEQ ID 26	gb AF046024 AF046024 Homo sapiens UBA3 (UBA3) mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF057569 AF057569 Homo sapiens upstream regulatory element binding protein 1 (UREB1) mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF065485 AF065485 Homo sapiens sorting nexin 4 mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF113615 AF113615 Homo sapiens FH1/FH2 domain-containing protein FHOS (FHOS) mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF117755 AF117755 Homo sapiens thyroid hormone receptor-associated protein complex component TRAP230 mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF132734 AF132734 Homo sapiens REC8 mRNA, partial cds.
Human OBRGRP	SEQ ID 26	gb AF141349 AF141349 Homo sapiens beta-tubulin mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF148213 AF148213 Homo sapiens aggrecanase-1 mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF161554 AF161554 Homo sapiens HSPC069 mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF195512 AF195512 Homo sapiens TNF2 (TNF2) mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF205588 AF205588 Homo sapiens ZNF01 and HUMORFKG1B genes, partial sequence, complete sequence.
Human OBRGRP	SEQ ID 26	gb AF205588 AF205588 Homo sapiens ZNF01 and HUMORFKG1B genes, partial sequence, complete sequence.
Human OBRGRP	SEQ ID 26	gb AF216493 AF216493 Homo sapiens a-helical protein (HCR) mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF224741 AF224741 Homo sapiens chloride channel protein 7 (CLCN7) mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AF233522 AF233522 Homo sapiens Golgi-associated, gamma-adaptin ear containing, ARF-binding protein 2 (GGA2) gene, complete cds.
Human OBRGRP	SEQ ID 26	gb AF279891 AF279891 Homo sapiens dead box protein 15 mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb AJ006267 HSAJ6267 Homo sapiens mRNA for ClpX-like protein.
Human OBRGRP	SEQ ID 26	gb AJ007798 HSA007798 Homo sapiens mRNA for stromal antigen 3 (STAG3 gene).
Human OBRGRP	SEQ ID 26	gb AJ131244 HSA131244 Homo sapiens mRNA for Sec24 protein (Sec24A isoform), partial.
Human OBRGRP	SEQ ID 26	gb AK001569 AK001569 Homo sapiens cDNA FLJ10707 fis, clone NT2RP3000859.
Human OBRGRP	SEQ ID 26	gb AK002174 AK002174 Homo sapiens cDNA FLJ11312 fis, clone PLACE1010105, weakly similar to RING CANAL PROTEIN.
Human OBRGRP	SEQ ID 26	gb AL031588 HS1163J1 Human DNA sequence from clone RP5-1163J1 on chromosome 22q13.2-13.33
Human OBRGRP	SEQ ID 26	gb AL117496 HSM801013 Homo sapiens mRNA; cDNA DKFZp434B0435 (from clone DKFZp434B0435); complete cds.

Human OBRGRP	SEQ ID 26	gb U40282 H SU40282 Homo sapiens integrin-linked kinase (ILK) mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb U58773 H SU58773 Human calcium binding protein mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb U59289 H SU59289 Human H-cadherin mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb U60325 H SU60325 Human DNA polymerase gamma mRNA, nuclear gene encoding mitochondrial protein, complete cds.
Human OBRGRP	SEQ ID 26	gb U65928 H SU65928 Human Jun activation domain binding protein mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb U78311 H SU78311 Human translation initiation factor 3 large subunit mRNA, complete cds.
Human OBRGRP	SEQ ID 26	gb X52022 H SCOLLV3 H.sapiens RNA for type VI collagen alpha3 chain.
Human OBRGRP	SEQ ID 26	gb X53416 H SABP280 Human mRNA for actin-binding protein (filamin) (ABP-280).
Human OBRGRP	SEQ ID 26	gb X80907 H SPHOSINK H.sapiens mRNA for p85 beta subunit of phosphatidylinositol-3-kinase.
Human OBRGRP	SEQ ID 26	gb X87241 H SHFATPRO H.sapiens mRNA for hFat protein.
Human OBRGRP	SEQ ID 26	gb Z49878 H SGACNMITS H.sapiens mRNA for guanidinoacetate N-methyltransferase.
Human Melatonin 1a receptor	SEQ ID 28	gb AB014522 AB014522 Homo sapiens mRNA for KIAA0622 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 28	gb AB018272 AB018272 Homo sapiens mRNA for KIAA0729 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 28	gb AC004797 AC004797 Homo sapiens chromosome 17, clone hRPC.62_O_9, complete sequence.
Human Melatonin 1a receptor	SEQ ID 28	gb AC011497 AC011497 Homo sapiens chromosome 19 clone CTB-50E14, complete sequence.
Human Melatonin 1a receptor	SEQ ID 28	gb AF156965 AF156965 Homo sapiens translocon-associated protein alpha subunit mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 28	gb AK000867 AK000867 Homo sapiens cDNA FLJ10005 fis, clone HEMBA1000156.
Human Melatonin 1a receptor	SEQ ID 28	gb AL157911 CNS01RGB Human chromosome 14 DNA sequence BAC R-16B13 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
Human Melatonin 1a receptor	SEQ ID 28	gb AQ210837 AQ210837 HS_2230_A1_D10_MR CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2230 Col=19 Row=G, DNA sequence.
Human Melatonin 1a receptor	SEQ ID 28	gb AQ636913 AQ636913 RPCI-11-465H9.TV RPCI-11 Homo sapiens genomic clone RPCI-11-465H9, DNA sequence.
Human Melatonin 1a receptor	SEQ ID 28	gb AQ893376 AQ893376 HS_3072_A1_D12_T7C CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=3072 Col=23 Row=G, DNA sequence.
Human Melatonin 1a receptor	SEQ ID 28	gb AR065358 AR065358 Sequence 2 from patent US 5849528.
Human Melatonin 1a receptor	SEQ ID 28	gb D63478 D63478 Human mRNA for KIAA0144 gene, complete cds.
Human Melatonin 1a receptor	SEQ ID 28	gb G30352 G30352 human STS SHGC-36823, sequence tagged site.

Human Melatonin 1a receptor	SEQ ID 28	gb L11316 MUSETX2X Mouse oncogene (ec2) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 28	gb M17886 HUMPPARP1 Human acidic ribosomal phosphoprotein P1 mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 28	gb M31166 HUMTSG14A Human tumor necrosis factor-inducible (TSG-14) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 28	gb U01062 HUMIP3R3 Human type 3 inositol 1,4,5-trisphosphate receptor (ITPR3) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB011148 AB011148 Homo sapiens mRNA for KIAA0576 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB014563 AB014563 Homo sapiens mRNA for KIAA0663 protein, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB020638 AB020638 Homo sapiens mRNA for KIAA0831 protein, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB020681 AB020681 Homo sapiens mRNA for KIAA0874 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB026190 AB026190 Homo sapiens mRNA for Kelch motif containing protein, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB028956 AB028956 Homo sapiens mRNA for KIAA1033 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB028981 AB028981 Homo sapiens mRNA for KIAA1058 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB032252 AB032252 Homo sapiens BAZ1A mRNA for bromodomain adjacent to zinc finger domain 1A, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB032253 AB032253 Homo sapiens BAZ1B mRNA for bromodomain adjacent to zinc finger domain 1B, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB035863 AB035863 Homo sapiens SCS-betaA mRNA for ATP specific succinyl CoA synthetase beta subunit precursor, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB037728 AB037728 Homo sapiens mRNA for KIAA1307 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AB037856 AB037856 Homo sapiens mRNA for KIAA1435 protein, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AC000118 HSAC000118 Human BAC clone RG072E11 from 7q21-7q22, complete sequence.
Human Melatonin 1a receptor	SEQ ID 30	gb AC005218 AC005218 Homo sapiens chromosome 5, P1 clone 737H5 (LBNL H36), complete sequence. L81819 L81820 L81821 L78765 AC002214 AC002215 AC000130
Human Melatonin 1a receptor	SEQ ID 30	gb AC005261 AC005261 Homo sapiens chromosome 19, CIT-HSP-444n24, complete sequence.
Human Melatonin 1a receptor	SEQ ID 30	gb AF007151 AF007151 Homo sapiens clone 23967 unknown mRNA, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF035191 AF035191 Homo sapiens nuclear autoantigenic sperm protein autosomal variant mRNA, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF044321 AF044321 Homo sapiens cytochrome c oxidase assembly protein COX11 (COX11) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF054284 AF054284 Homo sapiens spliceosomal protein SAP 155 mRNA, complete cds.

Human Melatonin 1a receptor	SEQ ID 30	gb AF058954 AF058954 Homo sapiens GTP-specific succinyl-CoA synthetase beta subunit (SCS) mRNA, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF084479 AF084479 Homo sapiens Williams-Beuren syndrome deletion transcript 9 (WBSCR9) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF086824 AF086824 Mus musculus rho/rac-interacting citron kinase (Crik) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF113699 AF113699 Homo sapiens clone FLB7527 PRO1999 mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF155116 AF155116 Homo sapiens NY-REN-60 antigen mRNA, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF161467 AF161467 Homo sapiens HSPC118 mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AF177198 AF177198 Homo sapiens talin mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AJ005197 HSAJ5197 Homo sapiens mRNA for guanine nucleotide exchange factor GRP1, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AK002174 AK002174 Homo sapiens cDNA FLJ11312 fis, clone PLACE1010105, weakly similar to RING CANAL PROTEIN.
Human Melatonin 1a receptor	SEQ ID 30	gb AL096791 HSJ659F15 Human DNA sequence from clone 659F15 on chromosome Xp11.21-11.4, complete sequence.
Human Melatonin 1a receptor	SEQ ID 30	gb AL137012 AL137012 Human DNA sequence from clone RP1-80G16 on chromosome 6, complete sequence.
Human Melatonin 1a receptor	SEQ ID 30	gb AL137714 HSM802200 Homo sapiens mRNA; cDNA DKFZp434K1323 (from clone DKFZp434K1323); partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb AL138995 CNS01DX0 Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC C-2588E21 of library CalTech-D from chromosome 14 of Homo sapiens (Human), complete sequence.
Human Melatonin 1a receptor	SEQ ID 30	gb AP000355 AP000355 Homo sapiens genomic DNA, chromosome 22q11.2, clone KB1896H10.
Human Melatonin 1a receptor	SEQ ID 30	gb D38521 HUMORF001 Human mRNA for KIAA0077 gene, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb D42039 HUMKIAAC Human mRNA for KIAA0081 gene, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb D44466 D44466 Homo sapiens mRNA for proteasome subunit p112, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb D78151 HUM26SPSP Human mRNA for 26S proteasome subunit p97, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb D87450 D87450 Human mRNA for KIAA0261 gene, partial cds.
Human Melatonin 1a receptor	SEQ ID 30	gb G24929 G24929 human STS EST204289, sequence tagged site.
Human Melatonin 1a receptor	SEQ ID 30	gb J04607 HUMTHRAA Human thyroid autoantigen mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb J39793 HUMNTF9 Homo sapiens nuclear factor p97 (NTF9) gene, complete cds.
Human Melatonin 1a receptor	SEQ ID 30	gb M16279 HUMMIC2A Human MIC2 mRNA, complete cds.

Human Melatonin 1a receptor	SEQ ID 32	gb AF134726 AF134726 Homo sapiens BAC clone 215O12 NG35, NG36, G9A, NG22, G9, HSP70-2, HSP70-1, HSP70-HOM, snRNP, G7A, NG37, NG23, and MutSH5 genes, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb AF156965 AF156965 Homo sapiens translocon-associated protein alpha subunit mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb AF162780 AF162780 Homo sapiens elastin microfibril interfase located protein (EMI) gene, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb AK000651 AK000651 Homo sapiens cDNA FLJ20644 fis, clone KAT02588.
Human Melatonin 1a receptor	SEQ ID 32	gb AQ636913 AQ636913 RPCI-11-465H9, TV RPCI-11 Homo sapiens genomic clone RPCI-11-465H9, DNA sequence.
Human Melatonin 1a receptor	SEQ ID 32	gb AQ773234 AQ773234 HS_2027_B1_A04_MR CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2027 Col=7 Row=B, DNA sequence.
Human Melatonin 1a receptor	SEQ ID 32	gb D14705 HUMALPHAC Human mRNA for alpha-catenin, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb D28476 HUMKGC1C Human mRNA for KIAA0045 gene, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb D42054 HUMKIAAM Human mRNA for KIAA0092 gene, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb G30352 G30352 human STS SHGC-36823, sequence tagged site.
Human Melatonin 1a receptor	SEQ ID 32	gb G50689 G50689 SHGC-83897 Human Homo sapiens STS genomic, sequence tagged site.
Human Melatonin 1a receptor	SEQ ID 32	gb J03210 HUMCN4GEL Human collagenase type IV mRNA, 3' end.
Human Melatonin 1a receptor	SEQ ID 32	gb L11316 MUSECT2X Mouse oncogene (ect2) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb L37418 HUME2K Homo sapiens dihydrolipoamide succinyltransferase (E2K) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb M33519 HUMBAT3A Human HLA-B-associated transcript 3 (BAT3) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb U12128 HSU12128 Human protein tyrosine phosphatase 1E (PTP1E) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb U17714 HSU17714 Homo sapiens putative tumor suppressor ST13 (ST13) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb U28918 HSU28918 Human progesterone receptor-associated p48 protein mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb U28964 HSU28964 Homo sapiens 14-3-3 protein mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb U47077 HSU47077 Homo sapiens DNA-dependent protein kinase catalytic subunit (DNA-PKcs) mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb U65928 HSU65928 Human Jun activation domain binding protein mRNA, complete cds.
Human Melatonin 1a receptor	SEQ ID 32	gb X02761 HFSIB1 Human mRNA for fibronectin (FN precursor).

Human Melatonin 1a receptor	SEQ ID 32	gb X15879 HSCOL1N Human mRNA for collagen VI alpha-1 N-terminal globular domain.
Human Melatonin 1a receptor	SEQ ID 32	gb X53416 HSABP280 Human mRNA for actin-binding protein (filamin) (ABP-280).
Human Melatonin 1a receptor	SEQ ID 32	gb X74008 HSPPPICC H.sapiens mRNA for protein phosphatase 1 gamma.
Human Melatonin 1a receptor	SEQ ID 32	gb Z97832 HS329A5 Human DNA sequence from clone RP3-329A5 on chromosome 6p21.1-21.33 Contains a pseudogene similar to ribosomal protein L35a, ZNF76 (zinc finger protein 76 (expressed in testis)), part of the gene for KIAA06460 protein, an EST, STSs, GSSs and CpG Islands.n, complete sequence.
Human melatonin 1b receptor	SEQ ID 34	gb AB014522 AB014522 Homo sapiens mRNA for KIAA0622 protein, partial cds.
Human melatonin 1b receptor	SEQ ID 34	gb AC004797 AC004797 Homo sapiens chromosome 17, clone hRPC:62_O_9, complete sequence.
Human melatonin 1b receptor	SEQ ID 34	gb AF064087 AF064087 Homo sapiens cullin 3 mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb AF086837 AF086837 Homo sapiens snapin mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb AF105253 AF105253 Homo sapiens neuroendocrine secretory protein 55 mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb AF177198 AF177198 Homo sapiens talin mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb AF188240 AF188240 Homo sapiens A1U mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb AF189009 AF189009 Homo sapiens ubiquitin-like product Chap1/Dsk2 mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb D28476 HUMKG1C Human mRNA for KIAA0045 gene, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb D79987 D79987 Homo sapiens mRNA for KIAA0165 gene, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb L27841 HUMPM1AUTO Human autoantigen pericentriol material 1 (PCM-1) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb L32602 RATOTX1X Rattus norvegicus OTX1 mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb L37418 HUME2K Homo sapiens dihydroliipoamide succinyltransferase (E2K) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb M15205 HUMTKRA Human thymidine kinase gene, complete cds, with clustered Alu repeats in the introns.
Human melatonin 1b receptor	SEQ ID 34	gb M81355 HUMSPHINO Homo sapiens sphingolipid activator proteins 1 and 2 processed mutant mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb U08815 HSU08815 Human splicesomal protein (SAP 61) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb U17714 HSU17714 Homo sapiens putative tumor suppressor ST13 (ST13) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 34	gb U28918 HSU28918 Human progesterone receptor-associated p48 protein mRNA,

Human melatonin 1b receptor	SEQ ID 36	gb M81355 HUMSPHINO Homo sapiens sphingolipid activator proteins 1 and 2 processed mutant mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U06863 HSU06863 Human follistatin-related protein precursor mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U17714 HSU17714 Homo sapiens putative tumor suppressor ST13 (ST13) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U33286 HSU33286 Human chromosome segregation gene homolog GAS mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U43368 HSU43368 Human VEGF related factor isoform VRF186 precursor (VRF) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U46025 HSU46025 Human translation initiation factor eIF-3 p110 subunit gene, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U47077 HSU47077 Homo sapiens DNA-dependent protein kinase catalytic subunit (DNA-PKcs) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U47077 HSU47077 Homo sapiens DNA-dependent protein kinase catalytic subunit (DNA-PKcs) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U59289 HSU59289 Human H-cadherin mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U65928 HSU65928 Human Jun activation domain binding protein mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U72398 HSU72398 Human Bcl-x beta (bcl-x) gene, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb U78311 HSU78311 Human translation initiation factor 3 large subunit mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 36	gb X51405 HSCARBE Human mRNA for carboxypeptidase E (EC 3.4.17.10).
Human melatonin 1b receptor	SEQ ID 36	gb Z11518 HSHRSRB H.sapiens mRNA for histidyl-tRNA synthetase.
Human melatonin 1b receptor	SEQ ID 38	gb AB007890 AB007890 Homo sapiens mRNA for KIAA0430 protein, partial cds.
Human melatonin 1b receptor	SEQ ID 38	gb AB018272 AB018272 Homo sapiens mRNA for KIAA0729 protein, partial cds.
Human melatonin 1b receptor	SEQ ID 38	gb AB023205 AB023205 Homo sapiens mRNA for KIAA0988 protein, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AB030653 AB030653 Homo sapiens mRNA for epsilon-adaptin, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AB032976 AB032976 Homo sapiens mRNA for KIAA1150 protein, partial cds.
Human melatonin 1b receptor	SEQ ID 38	gb AB037735 AB037735 Homo sapiens mRNA for KIAA1314 protein, partial cds.
Human melatonin 1b receptor	SEQ ID 38	gb AC007066 AC007066 Homo sapiens chromosome 9, clone hRPK.355_O_1, complete sequence.
Human melatonin 1b receptor	SEQ ID 38	gb AC010553 AC010553 Homo sapiens chromosome 16 clone RP11-59D8, complete sequence.
Human melatonin 1b receptor	SEQ ID 38	gb AF044321 AF044321 Homo sapiens cytochrome c oxidase assembly protein COX11 (COX11) mRNA, complete cds.

Human melatonin 1b receptor	SEQ ID 38	gb AF064087 AF064087 Homo sapiens cullin 3 mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF068755 AF068755 Homo sapiens sec7 domain family member (GBF1) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF072928 AF072928 Homo sapiens myotubularin related protein 6 mRNA, partial cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF086837 AF086837 Homo sapiens snapin mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF090397 QK17 Mus musculus QKI protein (qkl) gene, alternative splice products, exons 7a, 7b, and 7c and complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF113615 AF113615 Homo sapiens FH1/FH2 domain-containing protein FHOS (FHOS) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF119897 AF119897 Homo sapiens PRO2760 mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF189009 AF189009 Homo sapiens ubiquitin-like product Chap1/Dsk2 mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF208502 AF208502 Homo sapiens early B-cell transcription factor (EBF) mRNA, partial cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF216965 AF216965 Homo sapiens ancient conserved domain protein 3 (ACDP3) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AF229178 AF229178 Homo sapiens leucine rich repeat and death domain containing protein (LRDD) mRNA, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AJ005197 HSAJ5197 Homo sapiens mRNA for guanine nucleotide exchange factor GRP1, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb AK000331 AK000331 Homo sapiens cDNA FLJ20324 fis, clone HEP09841, highly similar to AB007931 Homo sapiens mRNA for KIAA0462 protein.
Human melatonin 1b receptor	SEQ ID 38	gb AK000331 AK000331 Homo sapiens cDNA FLJ20324 fis, clone HEP09841, highly similar to AB007931 Homo sapiens mRNA for KIAA0462 protein.
Human melatonin 1b receptor	SEQ ID 38	gb AK000867 AK000867 Homo sapiens cDNA FLJ10005 fis, clone HEMBA1000156.
Human melatonin 1b receptor	SEQ ID 38	gb AL136295 CNS01DVZ Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-468E2 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
Human melatonin 1b receptor	SEQ ID 38	gb AL137302 HSM801972 Homo sapiens mRNA; cDNA DKFZp434E146 (from clone DKFZp434E146).
Human melatonin 1b receptor	SEQ ID 38	gb D14705 HUMALPHAC Human mRNA for alpha-catenin, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb D28476 HUMKG1C Human mRNA for KIAA0045 gene, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb D38047 HUMPSP31 Human mRNA for 26S proteasome subunit p31, complete cds.
Human melatonin 1b receptor	SEQ ID 38	gb D50406 D50406 Homo sapiens ST15 mRNA, complete cds.

Human OB-receptor short form	SEQ ID 40	gb AF064087 AF064087 Homo sapiens cullin 3 mRNA, complete cds.
Human OB-receptor short form	SEQ ID 40	gb AK000867 AK000867 Homo sapiens cDNA FLJ10005 fis, clone HEMBA1000156.
Human OB-receptor short form	SEQ ID 40	gb AQ210837 AQ210837 HS_2230_A1_D10_MR CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2230 Col=19 Row=G, DNA sequence.
Human OB-receptor short form	SEQ ID 40	gb D42054 HUMKIAAM Human mRNA for KIAA0092 gene, complete cds.
Human OB-receptor short form	SEQ ID 40	gb G30352 G30352 human STS SHGC-36823, sequence tagged site.
Human OB-receptor short form	SEQ ID 40	gb L11316 MUSECT2X Mouse oncogene (ect2) mRNA, complete cds.
Human OB-receptor short form	SEQ ID 40	gb M32221 HUMSAPABCD Human saposin proteins A-D mRNA, complete cds.
Human OB-receptor short form	SEQ ID 40	gb M81355 HUMSPHINO Homo sapiens sphingolipid activator proteins 1 and 2 processed mutant mRNA, complete cds.
Human OB-receptor short form	SEQ ID 40	gb M81355 HUMSPHINO Homo sapiens sphingolipid activator proteins 1 and 2 processed mutant mRNA, complete cds.
Human OB-receptor short form	SEQ ID 40	gb U01062 HUMIP3R3 Human type 3 inositol 1,4,5-trisphosphate receptor (ITPR3) mRNA, complete cds.
Human OB-receptor short form	SEQ ID 40	gb U17714 HSU17714 Homo sapiens putative tumor suppressor ST13 (ST13) mRNA, complete cds.
Human OB-receptor short form	SEQ ID 40	gb U65928 HSU65928 Human Jun activation domain binding protein mRNA, complete cds.
hOB-receptor long form	SEQ ID 42	gb AB032254 AB032254 Homo sapiens BAZ2A mRNA for bromodomain adjacent to zinc finger domain 2A, complete cds.
hOB-receptor long form	SEQ ID 42	gb AC007537 AC007537 Homo sapiens 12p12 BAC RPC11-267J23 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
hOB-receptor long form	SEQ ID 42	gb AF003529 AF003529 Homo sapiens glypican 3 (GPC3) gene, partial cds and flanking repeat regions.
hOB-receptor long form	SEQ ID 42	gb AF157482 AF157482 Homo sapiens Rev7p (REV7) mRNA, complete cds.
hOB-receptor long form	SEQ ID 42	gb AF189009 AF189009 Homo sapiens ubiquitin-like product Chap1/Dsk2 mRNA, complete cds.
hOB-receptor long form	SEQ ID 42	gb AK000331 AK000331 Homo sapiens cDNA FLJ20324 fis, clone HEP09841, highly similar to AB007931 Homo sapiens mRNA for KIAA0462 protein.
hOB-receptor long form	SEQ ID 42	gb AK000867 AK000867 Homo sapiens cDNA FLJ10005 fis, clone HEMBA1000156.
hOB-receptor long form	SEQ ID 42	gb G30352 G30352 human STS SHGC-36823, sequence tagged site.
hOB-receptor long form	SEQ ID 42	gb L11316 MUSECT2X Mouse oncogene (ect2) mRNA, complete cds.
hOB-receptor long form	SEQ ID 42	gb M32221 HUMSAPABCD Human saposin proteins A-D mRNA, complete cds.
hOB-receptor long form	SEQ ID 42	gb M81355 HUMSPHINO Homo sapiens sphingolipid activator proteins 1 and 2 processed mutant mRNA, complete cds.

hOB-receptor long form	SEQ ID 42	gb U65928 HSU65928 Human Jun activation domain binding protein mRNA, complete cds.
hOB-receptor long form	SEQ ID 42	gb U74628 HSU74628 Homo sapiens cell division control related protein (hCDCrel-1) mRNA, complete cds.
hOB-receptor long form	SEQ ID 42	gb X74008 HSPPPICC H.sapiens mRNA for protein phosphatase 1 gamma.
hOB-receptor long form	SEQ ID 42	gb Z97832 HS329A5 Human DNA sequence from clone RP3-329A5 on chromosome 6p21.1-21.33 Contains a pseudogene similar to ribosomal protein L35a, ZNF76 (zinc finger protein 76 (expressed in testis)), part of the gene for KIAA06460 protein, an EST, STSs, GSSs and CpG Islands.n, complete sequence.
hOB-receptor long form	SEQ ID 44	gb AB032976 AB032976 Homo sapiens mRNA for KIAA1150 protein, partial cds.
hOB-receptor long form	SEQ ID 44	gb AC008958 AC008958 Homo sapiens chromosome 5 clone CTD-2353N24, complete sequence.
hOB-receptor long form	SEQ ID 44	gb AF064087 AF064087 Homo sapiens cullin 3 mRNA, complete cds.
hOB-receptor long form	SEQ ID 44	gb AF141327 AF141327 Homo sapiens ring finger protein BAP-1 mRNA, complete cds.
hOB-receptor long form	SEQ ID 44	gb AK000867 AK000867 Homo sapiens cDNA FLJ10005 fis, clone HEMBA1000156.
hOB-receptor long form	SEQ ID 44	gb AK001912 AK001912 Homo sapiens cDNA FLJ11050 fis, clone PLACE1004564, highly similar to CLEAVAGE AND POLYADENYLATION SPECIFICITY FACTOR, 100 KD SUBUNIT.
hOB-receptor long form	SEQ ID 44	gb AL135749 HSN14 Homo sapiens *** SEQUENCING IN PROGRESS *** from BAC CEPHB197N14, complete sequence.
hOB-receptor long form	SEQ ID 44	gb AL157911 CNS01RGB Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-16B13 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
hOB-receptor long form	SEQ ID 44	gb AL159199 HSPH21E4 H.sapiens STS from flow-sorted chromosome 13 random shear fragment, sequence tagged site.
hOB-receptor long form	SEQ ID 44	gb D42054 HUMKIAAM Human mRNA for KIAA0092 gene, complete cds.
hOB-receptor long form	SEQ ID 44	gb D63478 D63478 Human mRNA for KIAA0144 gene, complete cds.
hOB-receptor long form	SEQ ID 44	gb G30352 G30352 human STS SHGC-36823, sequence tagged site.
hOB-receptor long form	SEQ ID 44	gb L11316 MUSECT2X Mouse oncogene (ect2) mRNA, complete cds.
hOB-receptor long form	SEQ ID 44	gb M32886 HUMSRICPA Human sorcin CP-22 mRNA, complete cds.
hOB-receptor long form	SEQ ID 44	gb U70734 HSU70734 Homo sapiens 38 kDa Mov34 homolog mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb AC004770 AC004770 Homo sapiens chromosome 11, BAC CIT-HSP-311e8 (BC269730) containing the hFEN1 gene, complete sequence.
hOB-receptor long form	SEQ ID 46	gb AC004846 AC004846 Homo sapiens clone RP4-647C14, complete sequence.
hOB-receptor long form	SEQ ID 46	gb AC005517 AC005517 Homo sapiens chromosome 17, clone RP11-726O12, complete

hOB-receptor long form	SEQ ID 46	sequence. gb JAC005912 AC005912 Homo sapiens 12p13.3 BAC RPC11-543P15 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
hOB-receptor long form	SEQ ID 46	gb JAF006636 AF006636 Homo sapiens melanoma differentiation associated protein-9 (mda-9) mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb JAF086837 AF086837 Homo sapiens snapin mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb JAK000867 AK000867 Homo sapiens cDNA FLJ10005 fis, clone HEMBA1000156.
hOB-receptor long form	SEQ ID 46	gb AL157911 CNS01RGB Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-16B13 of library RPC1-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
hOB-receptor long form	SEQ ID 46	gb AL159199 HSPH21E4 H.sapiens STS from flow-sorted chromosome 13 random shear fragment, sequence tagged site.
hOB-receptor long form	SEQ ID 46	gb AL359611 HSM802730 Homo sapiens mRNA; cDNA DKFZp762L1710 (from clone DKFZp762L1710).
hOB-receptor long form	SEQ ID 46	gb AQ209770 AQ209770 HS_3244_A1_B01_T7 CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=3244 Col=1 Row=C, DNA sequence.
hOB-receptor long form	SEQ ID 46	gb AQ210837 AQ210837 HS_2230_A1_D10_MR CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2230 Col=19 Row=G, DNA sequence.
hOB-receptor long form	SEQ ID 46	gb D50063 HUMP40MOV Human mRNA for proteasome subunit p40 / Mov34 protein, complete cds.
hOB-receptor long form	SEQ ID 46	gb G24929 G24929 human STS EST204289, sequence tagged site.
hOB-receptor long form	SEQ ID 46	gb G30352 G30352 human STS SHGC-36823, sequence tagged site.
hOB-receptor long form	SEQ ID 46	gb L11316 MUSECT2X Mouse oncogene (ect2) mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb M17886 HUMPPARP1 Human acidic ribosomal phosphoprotein P1 mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb M32221 HUMSAPABCD Human saposin proteins A-D mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb M60119 HUMEP2AA Homo sapiens HIV-EP2/Schnurri-2 gene, complete cds.
hOB-receptor long form	SEQ ID 46	gb U17714 HSU17714 Homo sapiens putative tumor suppressor ST13 (ST13) mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb U28918 HSU28918 Human progesterone receptor-associated p48 protein mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb U47077 HSU47077 Homo sapiens DNA-dependent protein kinase catalytic subunit (DNA-PKcs) mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb U65928 HSU65928 Human Jun activation domain binding protein mRNA, complete cds.
hOB-receptor long form	SEQ ID 46	gb X78136 HSRNPE2 H.sapiens hnRNP-E2 mRNA.

Human ADBR kinase 1	SEQ ID 48	gb AB007949 AB007949 Homo sapiens mRNA for KIAA0480 protein, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AB008430 AB008430 Homo sapiens mRNA for CDEP, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AB011121 AB011121 Homo sapiens mRNA for KIAA0549 protein, partial cds.
Human ADBR kinase 1	SEQ ID 48	gb AB015617 AB015617 Homo sapiens ELKS mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AB017430 AB017430 Homo sapiens mRNA for kinesin-like DNA binding protein, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AB028956 AB028956 Homo sapiens mRNA for KIAA1033 protein, partial cds.
Human ADBR kinase 1	SEQ ID 48	gb AB033028 AB033028 Homo sapiens mRNA for KIAA1202 protein, partial cds.
Human ADBR kinase 1	SEQ ID 48	gb AB037825 AB037825 Homo sapiens mRNA for KIAA1404 protein, partial cds.
Human ADBR kinase 1	SEQ ID 48	gb AC023510 AC023510 Homo sapiens 12 BAC RP11-713N11 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
Human ADBR kinase 1	SEQ ID 48	gb AF031939 AF031939 Mus musculus RailBP1-associated EH domain protein Reps1 (reps1) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF084457 AF084457 Homo sapiens beta-cop homolog mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF056116 AF056116 Fugu rubripes serine/threonine kinase receptor type 1, All-1 related protein (ALR), fugu hedgehog (fhh), Ikaros-like, wnt1, wnt10b, ARF3, PAS1, and L41 ribosomal protein genes, complete cds; LRP1 gene, partial cds; and unknown genes.
Human ADBR kinase 1	SEQ ID 48	gb AF058718 AF058718 Homo sapiens putative 13 S Golgi transport complex 90kD subunit brain-specific isoform mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF084457 AF084457 Homo sapiens beta-cop homolog mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF124490 AF124490 Homo sapiens ARF GTPase-activating protein GIT1 mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF143946 AF143946 Homo sapiens transcriptional activator SRCAP (SRCAP) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF155135 AF155135 Homo sapiens novel retinal pigment epithelial cell protein (NORPEG) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF212162 AF212162 Homo sapiens ninein mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF216493 AF216493 Homo sapiens a-helical protein (HCR) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb AF231920 AF231920 Homo sapiens chromosome 21 unknown mRNA.
Human ADBR kinase 1	SEQ ID 48	gb AL096773 HS1000E10 Human DNA sequence from clone 1000E10 on chromosome 1p12-13.3, complete sequence.

Human ADBR kinase 1	SEQ ID 48	gb AL109658 HSJ776F14 Human DNA sequence from clone RP4-776F14 on chromosome 20p12.2-13. Contains the 5' end of the FKBP1A gene for FK506-binding protein 1A (12kD), the gene for the ortholog of mouse P47, part of the gene for a novel immunoglobulin domains containing protein, ESTs, STSs, GSSs and two putative CpG islands, complete sequence.
Human ADBR kinase 1	SEQ ID 48	gb AL139421 AL139421 Human DNA sequence from clone RP4-717I23 on chromosome 1p21.2-22.3, complete sequence.
Human ADBR kinase 1	SEQ ID 48	gb AL359235 CNS05TEM Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC C-3078G23 of library CalTech-D from chromosome 14 of Homo sapiens (Human), complete sequence.
Human ADBR kinase 1	SEQ ID 48	gb D17716 HUMNATV1 Human mRNA for N-acetylglucosaminyltransferase V, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb D21094 HUMHMP4 Human mRNA for motor protein, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb D87071 D87071 Human mRNA for KIAA0233 gene, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb G20945 G20945 human STS WI-30365, sequence tagged site.
Human ADBR kinase 1	SEQ ID 48	gb L04270 HUMTNFRFP Homo sapiens (clone CD18) tumor necrosis factor receptor 2 related protein mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb L04284 HUMHRX Human germline HRX mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb M13451 HUMLAMC Human lamin C mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb M32110 HUMP120PC Human proliferating-cell nuclear protein P120 mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb M80783 HUMB12A Human B12 protein mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb U10360 HSU10360 Human interferon-gamma gene, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb U12535 HSU12535 Human epidermal growth factor receptor kinase substrate (Eps8) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb U19348 HSU19348 Human (tpr-met fusion) oncogene mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb U36501 HSU36501 Human SP100-B (SP100-B) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb U50078 HSU50078 Human guanine nucleotide exchange factor p532 mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb U67963 HSU67963 Human lysophospholipase homolog (HU-K5) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb U85946 HSU85946 Homo sapiens brain secretory protein hSec10p (HSEC10) mRNA, complete cds.
Human ADBR kinase 1	SEQ ID 48	gb X57398 HSPM5 Human mRNA for pM5 protein.
Rat ADBR kinase 2	SEQ ID 50	gb AB002366 AB002366 Human mRNA for KIAA0368 gene, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB007931 AB007931 Homo sapiens mRNA for KIAA0462 protein, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB007963 AB007963 Homo sapiens mRNA for KIAA0494 protein, complete cds.

Rat ADBR kinase 2	SEQ ID 50	gb AB008430 AB008430 Homo sapiens mRNA for CDEP, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB015617 AB015617 Homo sapiens ELKS mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB017430 AB017430 Homo sapiens mRNA for kinesin-like DNA binding protein, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB020633 AB020633 Homo sapiens mRNA for KIAA0826 protein, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB020691 AB020691 Homo sapiens mRNA for KIAA0884 protein, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB028956 AB028956 Homo sapiens mRNA for KIAA1033 protein, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB029042 AB029042 Homo sapiens mRNA for ATPase inhibitor precursor, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB029331 AB029331 Homo sapiens HCR (a-helix coiled-coil rod homologue) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB033028 AB033028 Homo sapiens mRNA for KIAA1202 protein, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB037782 AB037782 Homo sapiens mRNA for KIAA1361 protein, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AB040951 AB040951 Homo sapiens mRNA for KIAA1518 protein, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AC005585 AC005585 Homo sapiens chromosome 22, clone hRPC.130_H_16, complete sequence.
Rat ADBR kinase 2	SEQ ID 50	gb AC008008 AC008008 Homo sapiens Xp22 PAC RPC16-102 (Roswell Park Cancer Institute Human PAC Library) complete sequence.
Rat ADBR kinase 2	SEQ ID 50	gb AF031463 AF031463 Homo sapiens phosphocin-like protein mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF042838 AF042838 Homo sapiens MEK kinase 1 (MEKK1) mRNA, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF044588 AF044588 Homo sapiens protein regulating cytokinesis 1 (PRC1) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF049105 AF049105 Homo sapiens centrosomal Nek2-associated protein 1 (C-NAP1) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF054284 AF054284 Homo sapiens spliceosomal protein SAP 155 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF058718 AF058718 Homo sapiens putative 13 S Golgi transport complex 90kD subunit brain-specific isoform mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF060181 AF060181 Homo sapiens zinc finger protein (ZNF198) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF063308 AF063308 Homo sapiens coiled-coil related protein DEEPEST (DEEPEST) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF064553 AF064553 Mus musculus NSD1 protein mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF067163 AF067163 Homo sapiens bamacan homolog mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF077599 AF077599 Homo sapiens hypothetical SBBI03 protein mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF083208 AF083208 Homo sapiens Che-1 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF084457 AF084457 Homo sapiens beta-cop homolog mRNA, complete cds.

Rat ADBR kinase 2	SEQ ID 50	gb AF100762 TRIP15 Homo sapiens thyroid receptor interactor trip15 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF100762 TRIP15 Homo sapiens thyroid receptor interactor trip15 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF113615 AF113615 Homo sapiens FH1/FH2 domain-containing protein FHOS (FHOS) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF131811 AF131811 Homo sapiens clone 24930 mRNA sequence.
Rat ADBR kinase 2	SEQ ID 50	gb AF143946 AF143946 Homo sapiens transcriptional activator SRCAP (SRCAP) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF161472 AF161472 Homo sapiens HSPC123 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF179308 AF179308 Homo sapiens KIF4 (KIF4) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF212162 AF212162 Homo sapiens ninein mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AF216493 AF216493 Homo sapiens a-helical protein (HCR) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb AJ001258 HSNIPSN1 Homo sapiens mRNA for NIPSNAP1 protein.
Rat ADBR kinase 2	SEQ ID 50	gb AJ005821 HSA5821 Homo sapiens mRNA for X-like 1 protein.
Rat ADBR kinase 2	SEQ ID 50	gb AK000160 AK000160 Homo sapiens cDNA FLJ20153 fis, clone COL08656, highly similar to AJ001381 Homo sapiens incomplete cDNA for a mutated allele.
Rat ADBR kinase 2	SEQ ID 50	gb AK001654 AK001654 Homo sapiens cDNA FLJ10792 fis, clone NT2RP4000560.
Rat ADBR kinase 2	SEQ ID 50	gb AK001858 AK001858 Homo sapiens cDNA FLJ10996 fis, clone PLACE1002433.
Rat ADBR kinase 2	SEQ ID 50	gb AK002174 AK002174 Homo sapiens cDNA FLJ11312 fis, clone PLACE1010105, weakly similar to RING CANAL PROTEIN.
Rat ADBR kinase 2	SEQ ID 50	gb AL050019 HSM800100 Homo sapiens mRNA; cDNA DKFZp564C186 (from clone DKFZp564C186).
Rat ADBR kinase 2	SEQ ID 50	gb AL162062 HSM802593 Homo sapiens mRNA; cDNA DKFZp762B245 (from clone DKFZp762B245); partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb AR035969 AR035969 Sequence 2 from patent US 5871970.
Rat ADBR kinase 2	SEQ ID 50	gb AZ095654 AZ095654 RPCI-23-476K9.TJ RPCI-23 Mus musculus genomic clone RPCI-23-476K9, DNA sequence.
Rat ADBR kinase 2	SEQ ID 50	gb D21094 HUMHMP4 Human mRNA for motor protein, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb D21260 HUMORFEA Human mRNA for KIAA0034 gene, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb D50063 HUMP40MOV Human mRNA for proteasome subunit p40 / Mov34 protein, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb D83777 D83777 Human mRNA for KIAA0193 gene, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb D87076 D87076 Human mRNA for KIAA0239 gene, partial cds.
Rat ADBR kinase 2	SEQ ID 50	gb G20945 G20945 human STS WI-30365, sequence tagged site.
Rat ADBR kinase 2	SEQ ID 50	gb J05243 HUMASPX Human nonerythroid alpha-spectrin (SPTAN1) mRNA, complete cds.

Rat ADBR kinase 2	SEQ ID 50	gb L04270 HUMTNFRRP Homo sapiens (clone CD18) tumor necrosis factor receptor 2 related protein mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb L12392 HUMHDA Homo sapiens Huntington's Disease (HD) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb L13848 HUMRNAHELA Human RNA helicase A mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb L20941 HUMFERRITH Human ferritin heavy chain mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb L27841 HUMPMIAUTO Human autoantigen pericentriol material 1 (PCM-1) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb L33075 HUMIQGA Homo sapiens ras GTPase-activating-like protein (IQGAP1) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb L39793 HUMNTF9 Homo sapiens nuclear factor p97 (NTF97) gene, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb L42572 HUMP8789R Homo sapiens p87/89 gene, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb M14144 HUMVIM Human vimentin gene, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb M32110 HUMP120PC Human proliferating-cell nucleolar protein P120 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb M55618 HUMHXB Homo sapiens hexabrachion (HXB) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb M62800 HUMSSARO Human 52-kD SS-A/Ro autoantigen mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb M80783 HUMB12A Human B12 protein mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb M84694 HUMMHHLAB4 Human MHC class I HLA-B*4005 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U01062 HUMIP3R3 Human type 3 inositol 1,4,5-trisphosphate receptor (ITPR3) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U03877 HSU03877 Human extracellular protein (S1-5) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U06863 HSU06863 Human follistatin-related protein precursor mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U10360 HSU10360 Human interferon-gamma gene, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U12535 HSU12535 Human epidermal growth factor receptor kinase substrate (Eps8) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U19348 HSU19348 Human (tpr-met fusion) oncogene mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U20285 HSU20285 Human Gps1 (GPS1) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U37139 HSU37139 Human beta 3-endonexin mRNA, long form and short form, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U39817 HSU39817 Human Bloom's syndrome protein (BLM) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U42390 HSU42390 Homo sapiens Trio mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U47077 HSU47077 Homo sapiens DNA-dependent protein kinase catalytic subunit (DNA-PKcs) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U52962 HSU52962 Human centrosomal protein kendrin mRNA, complete cds.

Rat ADBR kinase 2	SEQ ID 50	gb U53204 HSU53204 Human plectin (PLEC1) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U69668 HSU69668 Human nuclear pore complex-associated protein TPR (tpr) mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb U72761 HSU72761 Human karyopherin beta 3 mRNA, complete cds.
Rat ADBR kinase 2	SEQ ID 50	gb X13916 HSLDLRL Human mRNA for LDL-receptor related protein.
Rat ADBR kinase 2	SEQ ID 50	gb X15187 HSTRA1 Human tra1 mRNA for human homologue of murine tumor rejection antigen gp96.
Rat ADBR kinase 2	SEQ ID 50	gb X57398 HSPM5 Human mRNA for pM5 protein.
Rat ADBR kinase 2	SEQ ID 50	gb X91879 HSSPERMSP Homo sapiens mRNA for sperm protein.
Rat ADBR kinase 2	SEQ ID 50	gb X98801 HSDYNACTN H.sapiens mRNA for dynactin.
Rat ADBR kinase 2	SEQ ID 50	gb Z26649 HSPPLCB3 H.sapiens mRNA for phospholipase C-b3.
Rat ADBR kinase 2	SEQ ID 50	gb Z75330 HSSA1 H.sapiens mRNA for nuclear protein SA-1.
Rat ADBR kinase 2	SEQ ID 50	gb Z96932 HSP14PROT Homo sapiens mRNA for NA14 protein.
Rat beta Arrestin 1	SEQ ID 52	gb AB015856 AB015856 Homo sapiens mRNA for ATF6, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AB018541 AB018541 Homo sapiens PEX19 mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AB029290 AB029290 Homo sapiens mRNA for actin binding protein ABP620, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AB032254 AB032254 Homo sapiens BAZZA mRNA for bromodomain adjacent to zinc finger domain 2A, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AB041607 AB041607 Mus musculus brain cDNA, clone MNCb-5210, similar to Homo sapiens mRNA; cDNA DKFZp586B0519.
Rat beta Arrestin 1	SEQ ID 52	gb AC002369 AC002369 Homo sapiens Xp22 PAC RPC11-5G11 (from Roswell Park Cancer Center) complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AC004893 AC004893 Homo sapiens PAC clone RP4-808A1 from 7q21.1-q31.1, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AC007114 AC007114 Homo sapiens chromosome 17, clone hRPK.166_P_13, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AC007969 AC007969 Homo sapiens clone RP11-471A5, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AC009178 AC009178 Homo sapiens clone RP11-169C22 from 7q31, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AC009363 AC009363 Homo sapiens chromosome 14 clone CTD-2317F5 map 14q24.3, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AC009509 AC009509 Homo sapiens 12p11-37.2-54.4 BAC RP11-1060J15 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AC026694 AC026694 Homo sapiens chromosome 5 clone CTC-263A14, complete

Rat beta Arrestin 1	SEQ ID 52	sequence. gb AF000364 AF000364 Homo sapiens heterogeneous nuclear ribonucleoprotein R mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF016507 AF016507 Homo sapiens C-terminal binding protein 2 mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF035262 AF035262 Homo sapiens BAF57 (BAF57) gene, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF084940 AF084940 Homo sapiens beta-arrestin 1B mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF089841 AF089841 Homo sapiens gamma-filamin (ABPL) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF118073 AF118073 Homo sapiens PRO1748 mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF118082 AF118082 Homo sapiens PRO1902 mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF143946 AF143946 Homo sapiens transcriptional activator SRCAP (SRCAP) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF146692 AF146692 Homo sapiens filamin 2 (FLN2) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF170562 AF170562 Homo sapiens ubiquitin-specific processing protease (USP25) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF198097 AF198097 Homo sapiens chromosome Xp11.23 cosmid B167, E1017, and L2460, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AF233453 AF233453 Homo sapiens RACK-like protein PRKCBP1 (PRKCBP1) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AF279865 AF279865 Homo sapiens kinesin-like protein GAKIN mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb AL021579 HS598F2 Human DNA sequence from clone 598F2 on chromosome 1q23.1-24.3 Contains ESTs, STS and GSS, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AL080080 HSM800581 Homo sapiens mRNA; cDNA DKFZp564E1962 (from clone DKFZp564E1962); partial cds.
Rat beta Arrestin 1	SEQ ID 52	gb AL106427 CNS0168H Drosophila melanogaster genome survey sequence T7 end of BAC BACN15117 of DrosBAC library from Drosophila melanogaster (fruit fly), genomic survey sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AL109759 CNS0180W Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-898B23 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AL110204 HSM800856 Homo sapiens mRNA; cDNA DKFZp586K1922 (from clone DKFZp586K1922).
Rat beta Arrestin 1	SEQ ID 52	gb AL121576 CNS01DRW Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-476J6 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.

Rat beta Arrestin 1	SEQ ID 52	gb AL121733 HS126A53 Novel human gene mapping to chromosome 1.
Rat beta Arrestin 1	SEQ ID 52	gb AL133415 AL133415 Human DNA sequence from clone RP11-124N14 on chromosome 10. Contains the VIM gene for vimentin, the DNMT2 gene for DNA methyl transferase 2, the 5' end of the gene for intrinsic factor-B12 receptor precursor, ESTs, STSs, GSSs and two putative CpG islands, complete sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AL225216 CNS032RR Tetraodon nigroviridis genome survey sequence T7 end of clone 207F21 of library G from Tetraodon nigroviridis, genomic survey sequence.
Rat beta Arrestin 1	SEQ ID 52	gb AP000514 AP000514 Homo sapiens genomic DNA, chromosome 6p21.3, HLA Class I region, section 13/20.
Rat beta Arrestin 1	SEQ ID 52	gb AP001728 AP001728 Homo sapiens genomic DNA, chromosome 21q, section 72/105.
Rat beta Arrestin 1	SEQ ID 52	gb AQ694100 AQ694100 HS_2100_A2_G06_T7 CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2100 Col=12 Row=M, DNA sequence.
Rat beta Arrestin 1	SEQ ID 52	gb G22664 G22664 human STS WI-14136, sequence tagged site.
Rat beta Arrestin 1	SEQ ID 52	gb G24627 G24627 human STS WI-12266, sequence tagged site.
Rat beta Arrestin 1	SEQ ID 52	gb G30219 G30219 human STS SHGC-36528, sequence tagged site.
Rat beta Arrestin 1	SEQ ID 52	gb L39891 HUMPKD1GEN Homo sapiens polycystic kidney disease-associated protein (PKD1) gene, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb L78810 HUMYWWD703 Homo sapiens ADP/ATP carrier protein (ANT-2) gene, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb M58028 HUMUBIQAA Human ubiquitin-activating enzyme E1 (UBE1) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb M86667 HUMNAP H.sapiens NAP (nucleosome assembly protein) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb M99437 HUMNOTCH Human notch group protein (N) mRNA, partial cds.
Rat beta Arrestin 1	SEQ ID 52	gb U36600 HSU36600 Homo sapiens heparan N-deacetylase/N-sulfotransferase-1 mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb U77085 HSU77085 Human epithelial membrane protein (CL-20) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb U82761 HSU82761 Homo sapiens S-adenosyl homocysteine hydrolase homolog (XPV/kona) mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb U83867 HSU83867 Human alpha II spectrin mRNA, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb U88966 HSU88966 Human protein rapamycin associated protein (FRAP2) gene, complete cds.
Rat beta Arrestin 1	SEQ ID 52	gb X53416 HSABP280 Human mRNA for actin-binding protein (filamin) (ABP-280).
Rat beta Arrestin 1	SEQ ID 52	gb X57347 HSHS1RNA H.sapiens mRNA for HS1 protein.
Rat beta Arrestin 1	SEQ ID 52	gb X75315 HSRNASB4 H.sapiens seb4B mRNA.

Rat beta Arrestin 1	SEQ ID 52	gb X89984 HSBCL7A H.sapiens mRNA for BCL7A protein.
Rat beta aArrestin2	SEQ ID 54	gb AB002334 AB002334 Human mRNA for KIAA0336 gene, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AB007949 AB007949 Homo sapiens mRNA for KIAA0480 protein, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AB014600 AB014600 Homo sapiens mRNA for KIAA0700 protein, partial cds.
Rat beta aArrestin2	SEQ ID 54	gb AB018312 AB018312 Homo sapiens mRNA for KIAA0769 protein, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AB020716 AB020716 Homo sapiens mRNA for KIAA0909 protein, partial cds.
Rat beta aArrestin2	SEQ ID 54	gb AB022658 AB022658 Homo sapiens mRNA for KARP-1-binding protein 2 (KAB2), complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AB029002 AB029002 Homo sapiens mRNA for KIAA1079 protein, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AB037854 AB037854 Homo sapiens mRNA for KIAA1433 protein, partial cds.
Rat beta aArrestin2	SEQ ID 54	gb AC005297 AC005297 Homo sapiens Xp22-149 BAC GS1-466O4 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
Rat beta aArrestin2	SEQ ID 54	gb AC006372 AC006372 Homo sapiens clone RP11-331D5, complete sequence.
Rat beta aArrestin2	SEQ ID 54	gb AF068755 AF068755 Homo sapiens sec7 domain family member (GBF1) mRNA, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AF090170 AF090170 Homo sapiens Rad1-like protein (RAD1) mRNA, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AF126799 AF126799 Homo sapiens delta-6 fatty acid desaturase mRNA, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb AK000096 AK000096 Homo sapiens cDNA FLJ20089 fs, clone COL03992.
Rat beta aArrestin2	SEQ ID 54	gb AL021920 HS163M9 Homo sapiens DNA sequence from PAC 163M9 on chromosome 1p35.1-p36.21. Contains protein synthesis factor (eIF-4C), D1F15S1A pseudogene, ESTs, STS, GSS, complete sequence.
Rat beta aArrestin2	SEQ ID 54	gb AQ419619 AQ419619 RPCI-11-179P19.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-179P19, DNA sequence.
Rat beta aArrestin2	SEQ ID 54	gb AQ743015 AQ743015 HS_5387_B2_A10_T7A RPCI-11 Human Male BAC Library Homo sapiens genomic clone Plate=963 Col=20 Row=B, DNA sequence.
Rat beta aArrestin2	SEQ ID 54	gb AR063507 AR063507 Sequence 22 from patent US 5846711.
Rat beta aArrestin2	SEQ ID 54	gb I86850 I86850 Sequence 2 from patent US 5702903.
Rat beta aArrestin2	SEQ ID 54	gb L32832 HUMZFHP Homo sapiens zinc finger homeodomain protein (ATBF1-A) mRNA, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb M34175 HUMBADPTA Human beta adaptin mRNA, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb M61916 HUMLAM101 Human laminin B1 chain mRNA, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb M69043 HUMMAD3A Homo sapiens MAD-3 mRNA encoding IkB-like activity, complete cds.
Rat beta aArrestin2	SEQ ID 54	gb U26555 HSU26555 Human versican V2 core protein precursor splice-variant mRNA,

Rat beta aArestin2	SEQ ID 54	complete cds.
Rat beta aArestin2	SEQ ID 54	gb U35003 HSU35003 Human JNK2 beta2 protein kinase (JNK2B2) mRNA, complete cds.
Rat beta aArestin2	SEQ ID 54	gb U36188 HSU36188 Human clathrin assembly protein 50 (AP50) mRNA, complete cds.
Rat beta aArestin2	SEQ ID 54	gb U50078 HSU50078 Human guanine nucleotide exchange factor p532 mRNA, complete cds.
Rat beta aArestin2	SEQ ID 54	gb U73199 MMU73199 Mus musculus Rho-guanine nucleotide exchange factor mRNA, complete cds.
Rat beta aArestin2	SEQ ID 54	gb U79458 HSU79458 Human WW domain binding protein-2 mRNA, complete cds.
Rat beta aArestin2	SEQ ID 54	gb X13403 HSOCT1 Human mRNA for octamer-binding protein Oct-1.
Rat beta aArestin2	SEQ ID 54	gb X87241 HSHFATPRO H.sapiens mRNA for hFat protein.
Rat beta aArestin2	SEQ ID 54	gb Y13901 HSFGFR4G Homo sapiens FGFR-4 gene.
Rat beta aArestin2	SEQ ID 54	gb Z76735 HS246O8 Human DNA sequence from PAC 246O8, between markers DXS6791 and DXS8038 on chromosome X contains ESTs.
human STAT3	SEQ ID 56	gb AB002342 AB002342 Human mRNA for KIAA0344 gene, complete cds.
human STAT3	SEQ ID 56	gb AB006651 AB006651 Homo sapiens EXLM1 mRNA, complete cds.
human STAT3	SEQ ID 56	gb AB008430 AB008430 Homo sapiens mRNA for CDEP, complete cds.
human STAT3	SEQ ID 56	gb AB014537 AB014537 Homo sapiens mRNA for KIAA0637 protein, complete cds.
human STAT3	SEQ ID 56	gb AB018334 AB018334 Homo sapiens mRNA for KIAA0791 protein, complete cds.
human STAT3	SEQ ID 56	gb AB029012 AB029012 Homo sapiens mRNA for KIAA1089 protein, partial cds.
human STAT3	SEQ ID 56	gb AB033034 AB033034 Homo sapiens mRNA for KIAA1208 protein, partial cds.
human STAT3	SEQ ID 56	gb AC001644 AC001644 Genomic sequence from Human 9q34, complete sequence.
human STAT3	SEQ ID 56	gb AC005317 AC005317 Human Chromosome 15q26.1 PAC clone pDJ457j11 containing DNA polymerase gamma (polg) gene, complete sequence.
human STAT3	SEQ ID 56	gb AC008953 AC008953 Homo sapiens chromosome 5 clone CTD-2339M3, complete sequence.
human STAT3	SEQ ID 56	gb AC008997 AC008997 Homo sapiens chromosome 19 clone LLNLR-253D7, complete sequence.
human STAT3	SEQ ID 56	gb AF028832 AF028832 Homo sapiens Hsp89-alpha-delta-N mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF037439 AF037439 Homo sapiens protein kinase A anchoring protein mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF070656 AF070656 Homo sapiens FtsH homolog mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF105036 AF105036 Homo sapiens zinc finger transcription factor GKLF mRNA, complete cds.

human STAT3	SEQ ID 56	gb AF112207 AF112207 Homo sapiens translation initiation factor eIF-2b delta subunit mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF113615 AF113615 Homo sapiens FH1/FH2 domain-containing protein FHOS (FHOS) mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF153604 AF153604 Homo sapiens ubiquitin-specific protease homolog (UPH) mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF157476 AF157476 Homo sapiens DNA polymerase zeta catalytic subunit (REV3) mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF161541 AF161541 Homo sapiens HSPC056 mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF167173 AF167173 Homo sapiens chromosome X MSL3-1 protein mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF195951 AF195951 Homo sapiens signal recognition particle 68 mRNA, complete cds.
human STAT3	SEQ ID 56	gb AF197927 AF197927 Homo sapiens AF5q31 protein (AF5q31) mRNA, complete cds.
human STAT3	SEQ ID 56	gb AK000470 AK000470 Homo sapiens cDNA FLJ20463 fis, clone KAT06143.
human STAT3	SEQ ID 56	gb AK000703 AK000703 Homo sapiens cDNA FLJ20696 fis, clone KAI2488.
human STAT3	SEQ ID 56	gb AK001475 AK001475 Homo sapiens cDNA FLJ10613 fis, clone NT2RP2005393, weakly similar to AUTOANTIGEN NGP-1.
human STAT3	SEQ ID 56	gb AK001569 AK001569 Homo sapiens cDNA FLJ10707 fis, clone NT2RP3000859.
human STAT3	SEQ ID 56	gb AL162049 HSM802575 Homo sapiens mRNA; cDNA DKFZp762E1712 (from clone DKFZp762E1712); partial cds.
human STAT3	SEQ ID 56	gb D14705 HUMALPHAC Human mRNA for alpha-catenin, complete cds.
human STAT3	SEQ ID 56	gb D21260 HUMORFEA Human mRNA for KIAA0034 gene, complete cds.
human STAT3	SEQ ID 56	gb D28476 HUMKG1C Human mRNA for KIAA0045 gene, complete cds.
human STAT3	SEQ ID 56	gb D87077 D87077 Human mRNA for KIAA0240 gene, partial cds.
human STAT3	SEQ ID 56	gb G03796 G03796 human STS WI-1739.
human STAT3	SEQ ID 56	gb J03464 HUMC1A2 Human collagen alpha-2 type I mRNA, complete cds, clone pHCOL2A1.
human STAT3	SEQ ID 56	gb J03589 HUMUBILP Human ubiquitin-like protein (GdX) gene, complete cds.
human STAT3	SEQ ID 56	gb J03866 HUMIGMBC Homo sapiens M2 mitochondrial autoantigen dihydrolipoamide acetyltransferase mRNA, complete cds.
human STAT3	SEQ ID 56	gb L08069 HUMDNAJHOM Human heat shock protein, E. coli DnaJ homologue mRNA, complete cds.
human STAT3	SEQ ID 56	gb L21990 HUMSAP62X Human spliceosomal protein (SAP 62) gene, complete cds.
human STAT3	SEQ ID 56	gb L27841 HUMPM1AUTO Human autoantigen pericentriol material 1 (PCM-1) mRNA,

human STAT3	SEQ ID 56	complete cds.
human STAT3	SEQ ID 56	gb L29277 HUMAPRF Homo sapiens DNA-binding protein (APRF) mRNA, complete cds.
human STAT3	SEQ ID 56	gb L32832 HUMZFHP Homo sapiens zinc finger homeodomain protein (ATBF1-A) mRNA, complete cds.
human STAT3	SEQ ID 56	gb L47345 HUMELONA Homo sapiens elongin A mRNA, complete cds.
human STAT3	SEQ ID 56	gb M69066 HUMMOESIN Human moesin mRNA, complete cds.
human STAT3	SEQ ID 56	gb M77349 HUMTGFBIG Human transforming growth factor-beta induced gene product (BIGH3) mRNA, complete cds.
human STAT3	SEQ ID 56	gb M92439 HUM130LEU Human leucine-rich protein mRNA, complete cds.
human STAT3	SEQ ID 56	gb U12597 HSU12597 Human tumor necrosis factor type 2 receptor associated protein (TRAP3) mRNA, complete cds.
human STAT3	SEQ ID 56	gb U37122 HSU37122 Human adducin gamma subunit mRNA, complete cds.
human STAT3	SEQ ID 56	gb U39817 HSU39817 Human Bloom's syndrome protein (BLM) mRNA, complete cds.
human STAT3	SEQ ID 56	gb U41740 HSU41740 Human trans-Golgi p230 mRNA, complete cds.
human STAT3	SEQ ID 56	gb U42068 HSU42068 Human liver endoplasmic reticulum P58 mRNA, complete cds.
human STAT3	SEQ ID 56	gb U46571 HSU46571 Human tetrapeptide repeat protein (tpr2) mRNA, complete cds.
human STAT3	SEQ ID 56	gb U49056 RNU49056 Rattus norvegicus CTD-binding SR-like protein rA1 mRNA, complete cds.
human STAT3	SEQ ID 56	cds.
human STAT3	SEQ ID 56	gb U50078 HSU50078 Human guanine nucleotide exchange factor p532 mRNA, complete cds.
human STAT3	SEQ ID 56	gb U53204 HSU53204 Human plectin (PLEC1) mRNA, complete cds.
human STAT3	SEQ ID 56	gb U85245 HSU85245 Human phosphatidylinositol-4-phosphate 5-kinase type II beta mRNA, complete cds.
human STAT3	SEQ ID 56	gb X57398 HSPM5 Human mRNA for pM5 protein.
human STAT3	SEQ ID 56	gb X94910 HSERP28 Homo sapiens mRNA for ERp28 protein.
human STAT3	SEQ ID 56	gb X98801 HSDYNACTN H.sapiens mRNA for dynactin.
human STAT3	SEQ ID 56	gb Y09631 HSPIBF1 H.sapiens mRNA for PIBF1 protein, complete.
human STAT3	SEQ ID 56	gb Z68873 HSU209G1 Human DNA sequence from cosmid U209G1 on chromosome X.
human STAT3	SEQ ID 56	gb Z97184 HSF0811 Human DNA sequence from cosmid F0811 on chromosome 6. Contains Daxx, BING1, Tapasin, RGL2, KE2, BING4, BING5, ESTs and CpG islands.
human STAT3	SEQ ID 56	gb Z98044 HS510D11 Human DNA sequence from clone RP3-510D11 on chromosome 1p36.2-36.3
mCIS	SEQ ID 58	gb AB002451 AB002451 Homo sapiens mRNA from chromosome 5q21-22, clone A3-B.
mCIS	SEQ ID 58	gb AB011121 AB011121 Homo sapiens mRNA for KIAA0549 protein, partial cds.

mCIS	SEQ ID 58	gb AB020860 AB020860 Homo sapiens genomic DNA of 8p21.3-p22 anti-oncogene of hepatocellular colorectal and non-small cell lung cancer , segment 3/11.
mCIS	SEQ ID 58	gb AB026898 AB026898 Homo sapiens DNA, DLEC1 to ORCTL4 gene region, section 1/2 (DLEC1, ORCTL3, ORCTL4 genes, complete cds).
mCIS	SEQ ID 58	gb AB037856 AB037856 Homo sapiens mRNA for KIAA1435 protein, partial cds.
mCIS	SEQ ID 58	gb AC002551 HUAC002551 Human Chromosome 16 BAC clone CIT987SK-A-951C11, complete sequence.
mCIS	SEQ ID 58	gb AC004804 AC004804 Homo sapiens 12p13.3 PAC RPCI5-927J10 (Roswell Park Cancer Institute Human PAC library) complete sequence.
mCIS	SEQ ID 58	gb AC005480 AC005480 Homo sapiens clone DJ0592G07, complete sequence.
mCIS	SEQ ID 58	gb AC005913 AC005913 Homo sapiens Xp22-175-176 BAC GSHB-484O17 (Genome Systems Human BAC Library) complete sequence.
mCIS	SEQ ID 58	gb AC006238 AC006238 Homo sapiens chromosome 18, clone hRPK.474_N_24, complete sequence.
mCIS	SEQ ID 58	gb AC006518 AC006518 Homo sapiens 12p13 BAC RPCI11-144O23 (Roswell Park Cancer Institute Human BAC Library) complete sequence.
mCIS	SEQ ID 58	gb AC007263 AC007263 Homo sapiens chromosome 14 clone RP11-79J20 containing gene for checkpoint suppressor 1 (CHES1) gene, partial cds, complete sequence.
mCIS	SEQ ID 58	gb AC007272 AC007272 Homo sapiens clone RP11-13J8, complete sequence.
mCIS	SEQ ID 58	gb AC007934 AC007934 Homo sapiens, clone RP11-29A1, complete sequence.
mCIS	SEQ ID 58	gb AF000996 HSAF000996 Homo sapiens ubiquitous TPR motif, Y isoform (UTY) mRNA, alternative transcript 1, complete cds.
mCIS	SEQ ID 58	gb AF086837 AF086837 Homo sapiens snapin mRNA, complete cds.
mCIS	SEQ ID 58	gb AF090900 AF090900 Homo sapiens clone HQ0189 PRO0189 mRNA, complete cds.
mCIS	SEQ ID 58	gb AF146568 AF146568 Homo sapiens MIL 1 protein mRNA, complete cds; nuclear gene for mitochondrial product.
mCIS	SEQ ID 58	gb AF196779 AF196779 Homo sapiens transcription factor (GHM enhancer 3, JM11 protein, JM4 protein, JM5 protein, T54 protein, JM10 protein, A4 differentiation-dependent protein, triple LIM domain protein 6, and synaptophysin genes, complete cds; and L-type calcium channel alpha-1 subunit gene, partial cds, complete sequence.
mCIS	SEQ ID 58	gb AF196971 AF196971 Homo sapiens GATA-binding protein 1 and histone deacetylase-like protein genes, complete cds; CRAS pseudogene, complete sequence; and protein translocase gene, partial cds.
mCIS	SEQ ID 58	gb AF200348 AF200348 Homo sapiens melanoma-associated antigen MG50 mRNA, partial

mCIS		SEQ ID 58	cds. gb AF221842 AF221842 Homo sapiens U5 snRNP-associated 102 kDa protein mRNA, complete cds.
mCIS		SEQ ID 58	gb AJ001258 HSNIPSN1 Homo sapiens mRNA for NIPSNAP1 protein.
mCIS		SEQ ID 58	gb AK000331 AK000331 Homo sapiens cDNA FLJ20324 fis, clone HEP09841, highly similar to AB007931 Homo sapiens mRNA for KIAA0462 protein.
mCIS		SEQ ID 58	gb AL008734 HS324M8 Human DNA sequence from clone 324M8 on chromosome 1p36.2-36.3 Contains ESTs, GSSs and CpG Island, complete sequence.
mCIS		SEQ ID 58	gb AL021408 HS523C21 Homo sapiens DNA sequence from PAC 523C21 on chromosome 6q23.1-23.3. Contains EST, GSS and STS, complete sequence.
mCIS		SEQ ID 58	gb AL096857 HS598F21A Novel human mRNA from chromosome 1, which has similarities to BAT2 genes.
mCIS		SEQ ID 58	gb AL117355 CNS01DRL Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-354E14 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
mCIS		SEQ ID 58	gb AL133367 CNS01DUS Human chromosome 14 DNA sequence *** IN PROGRESS *** BAC R-600F24 of library RPCI-11 from chromosome 14 of Homo sapiens (Human), complete sequence.
mCIS		SEQ ID 58	gb AL136223 AL136223 Human DNA sequence from clone RP1-242G1 on chromosome 6p21.1-21.2, complete sequence.
mCIS		SEQ ID 58	gb AL137012 AL137012 Human DNA sequence from clone RP1-80G16 on chromosome 6, complete sequence.
mCIS		SEQ ID 58	gb AL359611 HSM802730 Homo sapiens mRNA; cDNA DKFZp762L1710 (from clone DKFZp762L1710).
mCIS		SEQ ID 58	gb AP001728 AP001728 Homo sapiens genomic DNA, chromosome 21q, section 72/105.
mCIS		SEQ ID 58	gb AQ395057 AQ395057 CITBI-E1-2542L3.TR CITBI-E1 Homo sapiens genomic clone 2542L3, DNA sequence.
mCIS		SEQ ID 58	gb AQ422416 AQ422416 RPCI-11-191D9.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-191D9, DNA sequence.
mCIS		SEQ ID 58	gb AQ568952 AQ568952 HS_5303_A1_C12_SP6E RPCI-11 Human Male BAC Library Homo sapiens genomic clone Plate=879 Col=23 Row=E, DNA sequence.
mCIS		SEQ ID 58	gb AQ609952 AQ609952 HS_5089_A1_F05_SP6E RPCI-11 Human Male BAC Library Homo sapiens genomic clone Plate=665 Col=9 Row=K, DNA sequence.

MCIS	SEQ ID 58	gb U85658 HSU85658 Human transcription factor ERF-1 mRNA, complete cds.
MCIS	SEQ ID 58	gb U91327 HSU91327 Human chromosome 12p15 BAC clone C1T987SK-99D8 complete sequence.
MCIS	SEQ ID 58	gb U94855 HSU94855 Homo sapiens translation initiation factor 3 47 kDa subunit mRNA, complete cds.
MCIS	SEQ ID 58	gb X52022 HSCOLLV3 H.sapiens RNA for type VI collagen alpha3 chain.
MCIS	SEQ ID 58	gb X54486 HSC1INHIB Human gene for C1-inhibitor.
MCIS	SEQ ID 58	gb Z68276 HSL190B4 Human DNA sequence from cosmid L190B4, Huntington's Disease Region, chromosome 4p16.3.
MCIS	SEQ ID 58	gb Z82195 HS274L7 Human DNA sequence from PAC 274L7 on chromosome X contains ESTs.
MCIS	SEQ ID 58	gb Z94044 HS154P24 Human DNA sequence from PAC 154P24 on chromosome X
MCIS	SEQ ID 58	gb Z99916 HS221G9 Human DNA sequence from clone CTA-221G9 on chromosome 22q11.21-12.2
MSOCS1	SEQ ID 60	gb AB037807 AB037807 Homo sapiens mRNA for KIAA1386 protein, partial cds.
MSOCS1	SEQ ID 60	gb AF064087 AF064087 Homo sapiens cullin 3 mRNA, complete cds.
MSOCS1	SEQ ID 60	gb AF161546 AF161546 Homo sapiens HSPC061 mRNA, complete cds.
MSOCS1	SEQ ID 60	gb J04543 HUMSNEXIN Human synexin mRNA, complete cds.
MSOCS1	SEQ ID 60	gb L13616 HUMFAKX Human focal adhesion kinase (FAK) mRNA, complete cds.
MSOCS1	SEQ ID 60	gb M96803 HUMSPTBN1A Human general beta-spectrin (SPTBN1) mRNA, complete cds.
MSOCS1	SEQ ID 60	gb U18543 HSU18543 Human zinc-finger protein mRNA, complete cds.
MSOCS1	SEQ ID 60	gb X14420 HSCOL3A1 Human mRNA for pro-alpha-1 type 3 collagen.